

LH LIQUID FORMULATIONS

Field of Invention

The invention relates to the field of liquid pharmaceutical formulations of luteinizing hormone (LH) as well as to methods of producing such formulations.

Background of the invention

Luteinizing hormone (LH), follicle-stimulating hormone (FSH) and chorionic gonadotropin (CG) are injectable proteins falling into the class of gonadotropins. LH, FSH and CG are used alone and in combination in the treatment of infertility and reproductive disorders in both female and male patients.

In nature, FSH and LH are produced by the pituitary gland. For pharmaceutical use, FSH and LH and their variants may be produced recombinantly (rFSH and rLH), or they may be produced from the urine of postmenopausal women (uFSH and uLH).

FSH is used in female patients in ovulation induction (OI) and in controlled ovarian hyperstimulation (COH) for assisted reproductive technologies (ART). In a typical treatment regimen for ovulation induction, a patient is administered daily injections of FSH or an FSH variant (about 75 to 300 IU FSH/day) for a period of from about 6 to about 12 days. In a typical treatment regimen for controlled ovarian hyperstimulation, a patient is administered daily injections of FSH or an FSH variant (about 150-600 IU FSH/day) for a period of from about 6 to about 12 days.

FSH is also used to induce spermatogenesis in men suffering from oligospermia. A regimen using 150 IU FSH 3 times weekly in combination with 2'500 IU hCG twice weekly has been successful in achieving an improvement in sperm count in men suffering from hypogonadotrophic hypogonadism (Burgues *et al.*, 1997).

LH is used in female patients in combination with FSH in OI and in COH, particularly in those patients having very low endogenous LH levels or resistance to LH, such as women suffering from hypogonadotrophic hypogonadism (HH, WHO group I) or older patients (i.e. 35 years or older), and patients in which embryo implantation or early miscarriage is a problem. LH in combination with FSH has traditionally been available in a preparation called human menopausal gonadotropins (hMG) extracted from the urine of postmenopausal women. hMG has a 1:1 ratio of FSH:LH activity.

CG acts at the same receptor as LH and elicits the same responses. CG has a longer circulation half-life than LH and is therefore commonly used as a long-acting source of LH-activity. CG is used in OI and COH regimens to mimic the natural LH peak and trigger ovulation. An injection of human chorionic gonadotropin (hCG) is used to trigger ovulation at the end of stimulation with FSH or a mixture of FSH and LH. CG may also be used together with FSH during stimulation for OI and COH, in order to provide LH-activity during stimulation in patients in which LH-activity is desirable, such as those mentioned above.

FSH, LH and CG are members of the heterodimer, glycoprotein hormone family that also includes thyroid stimulating hormone (TSH). The members of this family are heterodimers, comprising an α - and a β -subunit. The subunits are held together by noncovalent interactions. The human FSH (hFSH) heterodimer consists of (i) a mature 92 amino acid glycoprotein alpha subunit, which also is common to the other human family members (i.e., chorionic gonadotropin ("CG"), luteinizing hormone ("LH") and thyroid stimulating hormone ("TSH")); and (ii) a mature 111 amino acid beta subunit that is unique to FSH (Shome *et al.*, 1974 and 1988). The human LH heterodimer consists of (i) the mature 92 amino acid glycoprotein alpha subunit; and (ii) a mature 112 amino acid beta subunit that is unique to LH (Keutmann *et al.*, 1979; Talmadge *et al.*, 1984; Fiddes & Talmadge, 1984). The alpha and beta subunits of the glycoproteins may be prone to dissociate in formulations, due to interaction with a preservative, surfactant and other excipients. Dissociation of the subunits leads to loss of biological activity (Reichert & Ramsey, 1975).

FSH is formulated for intramuscular (IM) or subcutaneous (SC) injection. FSH is supplied in lyophilised (solid) form in vials or ampoules of 75 IU/vial and 150 IU/vial with a shelf life of about two years when stored at 2-25°C. A solution for injection is formed by reconstituting the lyophilised product with water for injection (WFI). For ovulation induction or controlled ovarian hyperstimulation, daily injections with starting doses of 75 IU to 600 IU are recommended for up to about ten days. Depending on the patient's response, up to three cycles of treatment with increasing doses of FSH can be used. With lyophilised formulations, the patient is required to reconstitute a new vial of lyophilised material with diluent and administer it immediately after reconstitution on a daily basis [Package insert N1700101A, published in February 1996, for Fertinex™ (urofollitropin for injection, purified) for subcutaneous injection, by Serono Laboratories, Inc., Randolph, MA].

FSH has also been formulated in both single-dose and multidose liquid formats, in vials, or ampoules. Single dose formats must remain stable and active in storage prior to use.

Multidose formats must not only remain stable and active in storage prior to use, but
 5 must also remain stable, active and relatively free of bacteria over the multiple day administration period, after the seal of the ampoule has been compromised. For this reason, multidose formats often contain a bacteriostatic agent.

LH is formulated for intramuscular (IM) or subcutaneous (SC) injection. LH is
 10 supplied in lyophilised (solid) form in vials or ampoules of 75 IU/vial with a shelf life of about two years when stored at 2-25°C. A solution for injection is formed by reconstituting the lyophilised product with water for injection (WFI). For ovulation induction or controlled ovarian hyperstimulation, in conjunction with FSH, daily injections with starting doses of 75 IU to 600 IU LH are recommended for up to about
 15 ten days.

EP 0 618 808 (Applied Research Systems ARS Holding N.V.) discloses a pharmaceutical composition comprising a solid intimate mixture of gonadotropin and a stabilising amount of sucrose alone or in combination with glycine.
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EP 0 814 841 (Applied Research Systems ARS Holding N.V.) discloses a stable, liquid pharmaceutical composition comprising recombinant human chorionic gonadotropin (hCG) and a stabilizing amount of mannitol.

25 WO 2004/087213 (Ares Trading S.A.) discloses a liquid and a freeze-dried pharmaceutical composition comprising follicle-stimulating hormone (FSH) or a variant thereof and/or luteinizing hormone (LH) or a variant thereof as well as a surfactant selected from Pluronic® F77, Pluronic F87, Pluronic F88 and Pluronic F68.

30 WO 2004/112826 (Ares Trading S.A.) discloses a freeze-dried formulation comprising follicle-stimulating hormone (FSH) or a variant thereof and luteinizing hormone (LH) or a variant thereof as well as a surfactant selected from a polysorbate including Tween 20, Tween 40 and Tween 80.

35 WO 00/04913 (Eli Lilly and Co.) discloses a formulation comprising FSH or an FSH variant, containing an alpha and beta subunit, and a preservative selected from the group consisting of phenol, m-cresol, p-cresol, o-cresol, chlorocresol, benzyl

alcohol, alkylparaben (methyl, ethyl, propyl, butyl and the like), benzalkonium chloride, benzethonium chloride, sodium dehydroacetate and thimerosal, or mixtures thereof in an aqueous diluent.

- 5 WO 2004/105788 (Ferring B.V.) discloses a pharmaceutical composition consisting of FSH and hCG in at least one pharmaceutically acceptable carrier.

EP 0 448 146 (AKZO N.V.) discloses a stabilized gonadotropin-containing lyophilisate comprising a gonadotropin and a stabilizing amount of a dicarboxylic acid
10 salt.

EP 0 853 945 (Akzo Nobel N.V.) discloses a liquid gonadotropin-containing formulation characterised in that the formulation comprises a gonadotropin and stabilising amounts of a polycarboxylic acid or a salt thereof and of a thioether
15 compound.

There remains a need for stable liquid formulations of LH or a LH variant either for single-dose or multidose administration.

20 **Summary of the invention**

It is an object of the invention to provide liquid formulations of LH or LH variant, methods for their preparation, and methods for their pharmaceutical or veterinary use in the treatment of fertility disorders.

- 25 In a first aspect, the invention provides a liquid formulation of LH or an LH variant comprising a purified preparation of LH or LH variant, a phosphate buffer and a stabilizing amount of arginine or salts thereof and/or lysine or salts thereof, according to claim 1.

- 30 In a second aspect, the invention provides a process for the preparation of a liquid formulation of LH or LH variant comprising the steps of a) forming a solution of LH or LH variant, and b) adding to said solution a stabilizing amount of arginine or salts thereof and/or lysine or salts thereof, and phosphate buffer, according to claim 23.

- 35 In a third aspect, the invention provides a presentation form of a liquid formulation of LH or LH variant comprising the liquid formulation according to the invention, hermetically sealed under sterile conditions in a container suitable for storage prior to

use, according to claim 22. The container may be an ampoule, a vial, a syringe, or a cartridge.

In a fourth aspect, the invention provides an article of manufacture for human pharmaceutical use, comprising packaging material and a container comprising the liquid formulation of LH or LH variant according to the invention and a bacteriostatic agent, wherein said packaging material comprises a label which indicates that such formulation may be held over a period of twenty-eight days or greater after the first use.

It is a further object of the invention to provide the use of a liquid formulation of LH or LH variant according to the invention in combination with liquid formulations of FSH or FSH variant, according to claim 26.

Brief description of the drawings

Figure 1 shows the comparison of chromatographic profiles (by SE-HPLC) of r-hLH formulations with different concentrations (6, 12 and 24 µg/ml of r-hLH), containers (syringe and cartridge) and filling volumes (0.25, 0.5 and 1 ml). These formulations, in addition to r-hLH, contain phosphate buffer, saccharose, Tween 20, methionine, benzyl alcohol, benzalkonium chloride and water for injection. The dotted lines correspond to a filling volume of 0.25 ml; the dashed lines correspond to a filling volume of 0.5 ml; and the black lines to a filling volume of 1 ml. (A) 6 µg/ml r-hLH formulation in syringe. (B) 6 µg/ml r-hLH formulation in cartridge. (C) 12 µg/ml r-hLH formulation in syringe. (D) 12 µg/ml r-hLH formulation in cartridge. (E) 24 µg/ml r-hLH formulation in syringe. (F) 24 µg/ml r-hLH formulation in cartridge. The peaks at a retention time of 8.5 minutes correspond to the intact heterodimers (alpha+beta subunits linked through a non-covalent bound), while peaks at higher retention time (about 9.5 minutes) correspond to the free-subunits. AU: absorbance unit.

Figure 2 shows the comparison of the percentage of heterodimers calculated by SE-HPLC on the formulation with and without the combination of bacteriostatic agents, respectively SAC/500/24µgBACL_cart and SAC/500/24µg_cart. (A) SE-HPLC performed after three days (3d) with storage at +40°C. (B) SE-HPLC performed after eight days (8d), four weeks (4w), six weeks (6w), eight weeks (8w) and thirteen weeks (13w) with storage at +33°C.

Detailed description of the invention

The liquid LH formulations of the present invention have improved or more suitable properties or stability, and are useful for infertility treatment in women and/or men.

5 In a preferred embodiment the liquid formulations of the invention are for subcutaneous and/or intramuscular injection.

The suitable properties or stability of the LH or LH variant liquid formulations according to the invention are obtained by preventing or reducing loss of activity or stability, or by improving any aspect of the effectiveness or desirability of
10 administration, e.g., by at least one of mode, frequency, dosage, comfort, ease of use, and the like.

Luteinising hormone, or LH, as used herein refers to the LH produced as a full-length mature protein, which includes, but is not limited to human LH or "hLH", whether
15 produced recombinantly or isolated from human sources, such as the urine of postmenopausal women.

The expression "LH variant" is meant to encompass those molecules differing in amino acid sequence, glycosylation pattern or in inter-subunit linkage from human LH
20 but exhibiting LH-activity.

The term "activity" in relation to LH, refers to the ability of an LH formulation or a mixed formulation, to elicit biological responses associated with LH, such as seminal vesicle weight gain method (Van Hell *et al.*, 1964). Biological activity of LH is
25 evaluated with respect to an accepted standard for LH.

Measure of activity is expressed in International Units per millilitre of solution (IU/ml) or in Mega International Units per millilitre of solution (MIU/ml). (1 MIU/ml = 1,000,000 IU/ml). An International Unit is calculated as described in the Research
30 Reference Reagent Note No. 35, published by the National Institute of Health, Bethesda, Maryland.

LH or LH variant can be produced by any suitable method, such as recombinantly, by isolation or purification from natural sources as may be the case, or by chemical
35 synthesis, or any combination thereof.

The use of the term "recombinant" refers to preparations of LH or a LH variant that are produced through the use of recombinant DNA technology (see for example WO 85/01958). One example of a method of expressing LH using recombinant technology is by transfection of eukaryotic cells with the DNA sequences encoding an alpha and beta subunit of LH, whether provided on one vector or on two vectors with each subunit having a separate promoter, as described in European patent nos. EP 0 211 894 and EP 0 487 512.

Another example of the use of recombinant technology to produce LH is by the use of homologous recombination to insert a heterologous regulatory segment in operative connection to endogenous sequences encoding the subunits of LH, as described in European patent no. EP 0 505 500 (Applied Research Systems ARS Holding NV).

Follicle stimulating hormone, or FSH, as used herein refers to the FSH produced as a full-length mature protein which includes, but is not limited to human FSH or "hFSH", whether produced recombinantly or isolated from human sources, such as the urine of postmenopausal women.

The expression "FSH variant" is meant to encompass those molecules differing in amino acid sequence, glycosylation pattern or in inter-subunit linkage from human FSH but exhibiting FSH-activity.

The term "activity" in relation to FSH, refers to the ability of an FSH formulation or a mixed formulation, to elicit biological responses associated with FSH, such as ovarian weight gain in the Steelman-Pohley assay (Steelman *et al.*, 1953), or follicular growth in a female patient. Follicular growth in a female patient can be evaluated by ultrasound, for example, in terms of the number of follicles having a mean diameter of at or about 16 mm on day 8 of stimulation. Biological activity is evaluated with respect to an accepted standard for FSH.

Examples of FSH variant include CTP-FSH, a long-acting modified recombinant FSH, consisting of the wild type α -subunit and a hybrid β -subunit in which the carboxy terminal peptide of hCG has been fused to the C-terminal of the β -subunit of FSH, as described in LaPolt *et al.* (1992) or in Klein *et al.* (2003). Also included is single chain CTP-FSH, a single chain molecule, consisting of the following sequences (from N-terminal to C-terminal):

β FSH	β hCG-CTP(113-145)	α FSH
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wherein β FSH signifies the β -subunit of FSH, β hCG CTP (113-145) signifies the
5 carboxy terminal peptide of hCG and α FSH signifies the α -subunit of FSH, as described by Klein *et al.* (2002).

Other examples of FSH variants include FSH molecules having additional glycosylation sites incorporated in the α - and/or β -subunit, as disclosed in WO 01/58493 (Maxygen), particularly as disclosed in claims 10 and 11 of WO 01/58493,
10 and FSH molecules with intersubunit S-S bonds, as disclosed in WO 98/58957.

FSH or FSH variant can be produced by any suitable method, such as recombinantly, by isolation or purification from natural sources as may be the case, or by chemical synthesis, or any combination thereof.

15 The FSH or FSH variant used in accordance with the present invention may be produced not only by recombinant means, including from mammalian cells, but also may be purified from other biological sources, such as from urinary sources. Acceptable methodologies include those described in Hakola *et al.* (1997), Keene *et al.* (1989), Cerpa-Poljak *et al.* (1993), Dias *et al.* (1994); Flack *et al.* (1994), Valove *et al.* (1994), U.S. Patent 3,119,740 and US Patent no. 5,767,067.
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The term "administer" or "administering" means to introduce a formulation of the present invention into the body of a patient in need thereof to treat a disease or
25 condition.

The term "patient" means a mammal that is treated for a disease or condition. Patients are of, but not limited to, the following origin, human, ovine, porcine, equine, bovine, rabbit and the like.

30 The term "aqueous diluent" refers to a liquid solvent that contains water. Aqueous solvent systems may be consist solely of water, or may consist of water plus one or more miscible solvents, and may contain dissolved solutes such as sugars, buffers, salts or other excipients. The more commonly used non-aqueous solvents are the short-chain organic alcohols, such as, methanol, ethanol, propanol, short-chain
35 ketones, such as acetone, and poly alcohols, such as glycerol.

An "isotonicity agent" is a compound that is physiologically tolerated and imparts a suitable tonicity to a formulation to prevent the net flow of water across cell membranes that are in contact with the formulation.

- 5 Compounds such as glycerin, are commonly used for such purposes at known concentrations. Other suitable isotonicity agents include, but are not limited to, amino acids or proteins (e.g., glycine or albumin), salts (e.g., sodium chloride), and sugars (e.g., dextrose, sucrose and lactose).
- 10 The term "bacteriostatic" or "bacteriostatic agent" refers to a compound or compositions added to a formulation to act as an anti-bacterial agent. A preserved LH or LH variant containing formulation of the present invention preferably meets statutory or regulatory guidelines for preservative effectiveness to be a commercially viable multi-use product, preferably in humans.
- 15 Examples of bacteriostatics include phenol, m-cresol, p-cresol, o-cresol, chlorocresol, benzyl alcohol, alkylparaben (methyl, ethyl, propyl, butyl and the like), benzalkonium chloride, benzethonium chloride, sodium dehydroacetate and thimerosal.

- 20 The term "buffer" or "physiologically-acceptable buffer" refers to solutions of compounds that are known to be safe for pharmaceutical or veterinary use in formulations and that have the effect of maintaining or controlling the pH of the formulation in the pH range desired for the formulation. Acceptable buffers for controlling pH at a moderately acidic pH to a moderately basic pH include, but are not limited to, such compounds as phosphate, acetate, citrate, arginine, TRIS, and
- 25 histidine. "TRIS" refers to 2-amino-2-hydroxymethyl-1,3,-propanediol, and to any pharmacologically acceptable salt thereof. Preferable buffers are phosphate buffers with saline or an acceptable salt.

- 30 The term "phosphate buffer" refers to solutions containing phosphoric acid or salts thereof, adjusted to a desired pH. Generally phosphate buffers are prepared from phosphoric acid, or a salt of phosphoric acid, including but not limited to sodium and potassium salts. Several salts of phosphoric acid are known in the art, such as sodium and potassium monobasic, dibasic, and tribasic salts of the acid. Salts of phosphoric acid are also known to occur as hydrates of the occurring salt. Phosphate
- 35 buffers may cover a range of pHs, such as from about pH 4 to about pH 10, and preferred ranges from about pH 5 to about pH 9, and a most preferred range of at or about 7.5 to at or about 8.5, most preferably at or about pH 8.0.

The term "vial" or "container" refers broadly to a reservoir suitable for retaining LH and diluent in a contained sterile state. Examples of a vial as used herein include ampoules, cartridges, blister packages, or other such reservoir suitable for delivery of the LH to the patient via syringe, pump (including osmotic), catheter, transdermal patch, pulmonary or transmucosal spray. Vials suitable for packaging products for parenteral, pulmonary, transmucosal, or transdermal administration are well known and recognized in the art.

The term "stability" refers to the physical, chemical, and conformational stability of LH in the formulations of the present invention (including maintenance of biological activity). Instability of a protein formulation may be caused by chemical degradation or aggregation of the protein molecules to form higher order polymers, by dissociation of the heterodimers into monomers, deglycosylation, modification of glycosylation, oxidation (particularly of the α -subunit) or any other structural modification that reduces at least one biological activity of an LH polypeptide included in the present invention.

A "stable" solution or formulation, is one wherein the degree of degradation, modification, aggregation, loss of biological activity and the like, of proteins therein is acceptably controlled, and does not increase unacceptably with time. Preferably the formulation retains at least at or about 80% of the labelled LH activity over a period of 6 months at a temperature of at or about 1-10°C, more preferably at or about 2-8°C, more preferably at or about 4-5°C. LH activity can be measured using the seminal vesicle weight gain bioassay⁶.

The term "treating" refers to the administration, follow up, management and/or care of a patient for which LH administration is desirable for the purpose of follicle or testicular stimulation or any other physiological response regulated by LH. Treating can thus include, but is not limited to, the administration of LH for the induction or improvement of sperm quality, stimulation of testosterone release in the male, or follicular development or for ovulation induction in the female.

The expression "multidose use" is intended to include the use of a single ampoule, vial, or cartridge of an LH formulation for more than one injection, for example 2, 3, 4, 5, 6 or more injections. The injections are preferably made over a period of at least at or about 12 hours, 24 hours, 48 hours, etc., preferably up to a period of at or about 28 days.

The injections may be spaced in time, for example, by a period of 6, 12, 24, 48 or 72 hours.

5 A "salt" of a protein is an acid or base addition salt. Such salts are preferably formed between any one or more of the charged groups in the protein and any one or more physiologically acceptable, non-toxic cations or anions. Organic and inorganic salts include, for example, those prepared from acids such as hydrochloric, sulphuric, sulfonic, tartaric, fumaric, hydrobromic, glycolic, citric, maleic, phosphoric, succinic, acetic, nitric, benzoic, ascorbic, p-toluenesulfonic, benzenesulfonic,
10 naphthalenesulfonic, propionic, carbonic, and the like, or for example, ammonium, sodium, potassium, calcium, or magnesium.

It has been found that an amino acid selected from the group of arginine, lysine or a mixture or a salt thereof is a suitable stabilizing agent for preparing a stable liquid
15 formulation comprising LH. Therefore, the first aspect of the invention relates to a liquid gonadotropin formulation containing luteinizing hormone (LH) or a variant thereof characterized in that said formulation comprises a stabilizing amount of arginine or salts thereof and/or lysine or salts thereof.

20 The concentration of arginine or salts thereof is at 10 mg/ml to 50 mg/ml, more preferably at or about 20 mg/ml to at or about 40 mg/ml, more particularly preferably at or about 25 mg/ml to at or about 35 mg/ml, most preferably at or about 31.5 mg/ml.

The concentration of lysine or salts thereof is at 10 mg/ml to 50 mg/ml, more
25 preferably at or about 20 mg/ml to at or about 40 mg/ml, more particularly preferably at or about 25 mg/ml to at or about 35 mg/ml, most preferably at or about 28.5 mg/ml.

The luteinizing hormone (LH) within the liquid formulation is preferably present at a concentration of at or about 1 to at or about 50 µg/ml of the total formulation.
30 In one embodiment, the luteinizing hormone (LH) is present at a concentration of at or about 1 to 15 µg/ml of the total formulation, in particular when intended for single use.

In a further embodiment, the luteinizing hormone (LH) is present at a concentration of
35 at or about 15 to 30 µg/ml of the total formulation, in particular when intended for multiple use (multidose).

The LH concentration in the formulation is preferably at or about 20 IU/ml to at or about 2,000 IU/ml, more preferably at or about 50 to at or about 1,000 IU/ml, more particularly preferably at or about 100 IU/ml to at or about 600 IU/ml.

- 5 Preferably the LH is produced recombinantly, particularly preferably it is produced in Chinese hamster ovary (CHO) cells transfected with an expression vector or vectors comprising DNA coding for the human glycoprotein alpha-subunit and the beta-subunit of LH. DNA encoding the alpha and beta-subunits may be present on the same or different vectors.

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Recombinant LH has several advantages over its urinary counterparts. Culture and isolation techniques using recombinant cells permit consistency between batches. In contrast, urinary LH varies greatly from batch to batch in such characteristics as purity, glycosylation pattern, sialylation and oxidation of the subunits. Due to greater
 15 batch-to-batch consistency and purity of recombinant LH, the hormones can be readily identified and quantified using techniques such as isoelectric focusing (IEF). The ease with which recombinant LH can be identified and quantified permits the filling of vials by mass of hormone (fill-by-mass) rather than filling by bioassay.

- 20 Preferably the liquid formulations of the present invention have a buffer, preferably a phosphate buffer, with preferred counterions being sodium or potassium ions. Phosphate saline buffers are well known in the art, such as Dulbecco's Phosphate buffered saline. Buffer concentrations in total solution can vary between at or about 1mM, 5mM, 9.5mM, 10mM, 50mM, 100mM, 150mM, 200mM, 250mM, and 500mM.
 25 Preferably the buffer concentration is at or about 10mM. Particularly preferred is a buffer 10 mM in phosphate ions with a pH of at or about 8.0.

- Preferably the buffer is adjusted in such a way that the liquid formulations of the present invention have a pH between at or about 7.0 and at or about 9.0, more
 30 preferably at or about 7.5 to at or about 8.5, including about pH 7.8, pH 8.0, and 8.2. The invention is directed to liquid formulations of LH or LH variant that may be single dose or multidose.

- Those liquid LH formulations of the invention that are intended for multidose use
 35 comprise a bacteriostatic agent, such as phenol, m-cresol, p-cresol, o-cresol, chlorocresol, benzyl alcohol, alkylparaben (methyl, ethyl, propyl, butyl and the like), benzalkonium chloride, benzethonium chloride, sodium dehydroacetate and

thimerosal. Preferred are benzyl alcohol, phenol, and combination of benzyl alcohol and benzalkonium chloride. The bacteriostatic agent is used in an amount that will yield a concentration that is effective to maintain the formulation essentially bacteria free (suitable for injection) over the multidose injection period, which may be at or
 5 about 12 or 24 hours to at or about 12 or 14 days, preferably at or about 6 to at or about 28 days.

The bacteriostatic agent is preferably present in a concentration of at or about 0.005 to at or about 15 mg/ml, more preferably at or about 0.01 to at or about 12 mg/ml.
 10 The bacteriostatic agent is preferably present in a concentration of at or about 0.1% (mass bacteriostatic/mass of solvent) to at or about 2.0%, more preferably at or about 0.2% to at or about 1.0%.

In the case of benzyl alcohol, particularly preferred is a concentration of 0.9% or 1.2%. In the case of phenol, particularly preferred is at or about 0.5%. In the case of
 15 combination of benzyl alcohol and benzalkonium chloride, particularly preferred is a concentration of 0.3% and 0.001%, respectively.

Preferably the formulations of the invention contain an antioxidant, such as methionine, sodium bisulfite, salts of ethylenediaminetetraacetic acid (EDTA),
 20 butylated hydroxytoluene (BHT), and butylated hydroxy anisole (BHA). Most preferred is methionine. The antioxidant prevents oxidation of LH (particularly the α -subunit).

The antioxidant, e.g. methionine is preferably present at a concentration of at or
 25 about 0.01 to at or about 5.0 mg/ml, more preferably at or about 0.05 to at or about 0.5 mg/ml.

Preferably the formulations of the invention contain a surfactant. Preferably the surfactant is selected from the group of polysorbates, in particular Tween 20
 30 (polyoxyethylene (20) sorbitan monolaurate), Tween 40 (polyoxyethylene (20) sorbitan monopalmitate) and Tween 80 (polyoxyethylene (20) sorbitan monooleate). Most preferred is Tween 20, preferably at a concentration of at or about 0.01 to at or about 10 mg/ml.

35 In a preferred embodiment the formulations of the invention comprise at least 25 μ g/ml of r-hLH, 1.65 mg/ml of $\text{Na}_2\text{HPO}_4 \cdot 2\text{H}_2\text{O}$, 0.104 mg/ml of $\text{NaH}_2\text{PO}_4 \cdot \text{H}_2\text{O}$, 31.5

mg/ml of arginine monohydrochloride, 0.05 mg/ml of Tween 20, 0.5 mg/ml of methionine and 5 mg/ml of phenol.

5 In another preferred embodiment the formulations of the invention comprise at least 25 µg/ml of r-hLH, 1.65 mg/ml of Na₂HPO₄ 2H₂O, 0.104 mg/ml of NaH₂PO₄ H₂O, 28.5 mg/ml of lysine monohydrochloride, 0.05 mg/ml of Tween 20, 0.5 mg/ml of methionine and 12 mg/ml of benzylalcohol.

10 In another preferred embodiment the formulations of the invention comprise at least 25 µg/ml of r-hLH, 1.65 mg/ml of Na₂HPO₄ 2H₂O, 0.104 mg/ml of NaH₂PO₄ H₂O, 28.5 mg/ml of lysine monohydrochloride, 0.05 mg/ml of Tween 20, 0.5 mg/ml of methionine 3 mg/ml of benzylalcohol and 0.01 mg/ml of benzalkonium chloride.

15 In a preferred embodiment, the invention provides a liquid pharmaceutical composition, for multidose use, comprising LH or LH variant, a stabilizing agent selected from the group of arginine, lysine or a mixture or a salt thereof, and a bacteriostatic agent selected from benzyl alcohol, phenol, and a combination of benzyl alcohol and benzalkonium chloride.

20 In a further preferred embodiment, the invention provides a method for manufacturing a liquid pharmaceutical composition, for multidose use, comprising forming an aqueous solution of LH or LH variant, a stabilizing agent selected from the group of arginine, lysine or a mixture or a salt thereof, and a bacteriostatic agent selected from benzyl alcohol, phenol, and a combination of benzyl alcohol and benzalkonium
25 chloride, and WFI (water for injection).

In yet another preferred embodiment, the invention provides a method for manufacturing a packaged pharmaceutical composition comprising dispensing a solution comprising LH or LH variant, a stabilizing agent selected from the group of
30 arginine, lysine or a mixture or a salt thereof, and a bacteriostatic agent selected from benzyl alcohol, phenol, and a combination of benzyl alcohol and benzalkonium chloride.

In yet another preferred embodiment, the invention provides an article of
35 manufacture for human pharmaceutical use, comprising a vial comprising a solution of LH or LH variant, a stabilizing agent selected from the group of arginine, lysine or

a mixture or a salt thereof, and a bacteriostatic agent selected from benzyl alcohol, phenol, and a combination of benzyl alcohol and benzalkonium chloride, and a label stating that such solution may be held over a period of at or about twenty-four hours or greater after the first use. Preferably the label states that the solution may be held up to at or about 12 or 14 days after the first use.

Before the first use, that is before the seal of the vial ampoule or cartridge has been broken, the formulations of the invention may be kept for at least at or about 6 months, 12 months or 24 months.

10 Under preferred storage conditions, before the first use, the formulations are kept away from bright light (preferably in the dark), at temperatures of at or about 2-8°C, more preferably at or about 4-5°C.

As noted above, the invention provides liquid formulations of LH or LH variant for single use and multidose use, containing a bacteriostatic. The formulations of the invention are suitable for pharmaceutical or veterinary use.

As noted above, in a preferred embodiment, the invention provides an article of manufacture, comprising packaging material and a vial comprising a solution of LH or LH variant, a stabilizing agent selected from the group of arginine, lysine or a mixture or a salt thereof, a bacteriostatic agent selected from benzyl alcohol, phenol, and a combination of benzyl alcohol and benzalkonium chloride, optionally with buffers and/or other excipients, in an aqueous diluent, wherein said packaging material comprises a label which indicates that such solution may be held over a period of twenty-four hours or greater after the first use.

Preferably the formulations of the invention retain at least at or about 80% of the LH activity at the time of packaging over a period of 24 months (before the first use). LH activity can be measured using the rat seminal vesicle weight gain bioassay⁵.

30 The formulations of the present invention can be prepared by a process which comprises mixing LH or a variant thereof and a stabilizing amount of arginine or salts thereof and/or lysine or salts thereof, and optionally a bacteriostatic agent selected from benzyl alcohol, phenol, and a combination of benzyl alcohol and benzalkonium chloride as solids or dissolving LH or a variant thereof and a stabilizing amount of arginine or salts thereof and/or lysine or salts thereof, and optionally a bacteriostatic agent selected from benzyl alcohol, phenol, and a combination of benzyl alcohol and

benzalkonium chloride in an aqueous diluent. Mixing the components and dissolving them in an aqueous diluent is carried out using conventional dissolution and mixing procedures. To prepare a suitable formulation, for example, a measured amount of LH or LH variant in buffered solution is combined with arginine or salts thereof and/or lysine or salts thereof and optionally a bacteriostatic selected from benzyl alcohol, phenol, and a combination of benzyl alcohol and benzalkonium chloride in a buffered solution in quantities sufficient to provide the protein, arginine or salts thereof and/or lysine or salts thereof and the optional bacteriostatic at the desired concentrations. The resulting solution is then dispensed into vials, ampoules or cartridges. Variations of this process would be recognized by one of ordinary skill in the art.

For example, the order the components are added, whether additional additives are used, the temperature and pH at which the formulation is prepared, are all factors that may be optimised for the concentration and means of administration used.

In a preferred embodiment, the formulations of the invention are made by preparing individual stock solutions of known concentration of all the components of the formulation (e.g. buffer sodium phosphate, arginine or salts thereof and/or lysine or salts thereof, Tween 20, methionine, LH), and aliquoting volumetric amounts to form a “mother solution” of the same composition as the final formulation. The “mother solution” is preferably filtered through a Duropore[®] (Millipore) 0.22 micron PDF membrane, to remove microorganisms, and then aliquots are dispensed into individual containers, such as vials, ampoules or cartridges.

The formulations of the present invention can be used in combination with formulations comprising FSH or FSH variant (e.g. Gonal-F[®]).

The formulations of the present invention can be administered using recognized devices. Examples comprising these single vial systems include pen-injector devices for delivery of a solution such as EasyJect[®], Gonal-F[®] Pen, Humaject[®], NovoPen[®], B-D[®] Pen, AutoPen[®], and OptiPen[®].

The products presently claimed include packaging material. The packaging material provides, in addition to the information required by the regulatory agencies, the conditions under which the product may be used. For the single vial, solution product, the label indicates that such solution may be stored after first use for a period of

twenty-four hours or greater, preferably for up to 12 or 14 days. The presently claimed products are useful for human pharmaceutical product use.

5 The following examples are provided merely to further illustrate the preparation of the formulations and compositions of the invention. The scope of the invention shall not be construed as merely consisting of the following examples.

Examples

Materials

Item	Manufacturer
r-hLH Bulk	Laboratoires Serono SA
L-Arginine monohydrochloride	Merck code 1.01544
Benzalkonium chloride	Fluka code 12063
Benzyl alcohol	Merck code 1.00987
m-Cresol	Merck code 8.09691
L-Glycine	Merck code 5.00190
L-Lysine monohydrochloride	Merck code 1.05701
Methionine	Merck code 1.05707
Phenol	Merck code 1.00200
Ortho-phosphoric Acid 85% (Ph Eur, BP, NF)	Merck code 1.00563
Sodium hydroxide	Merck code 1.06498
di-Sodium hydrogen phosphate dihydrate	Merck code 1.06580
Sodium dihydrogen phosphate monohydrate	Merck code 1.06346
Sodium sulphate anhydrous	Merck code 1.006649
Sorbitol	Sigma code S-1876
Saccharose	Merck code 1.07653
Trehalose	Merck code 1.08216
Tween 20	Merck code 8.22184
Water	milliQ

5 The following study evaluated the following parameters for a large number of formulations:

- Investigation on stabilizing and/or bacteriostatic agents
- Compatibility with FSH formulations
- Compatibility with primary packaging

10 • Stability profile after storage

The formulations were liquid monodose and multidose formulations. The following six stabilizing agents were evaluated:

- L-Arginine monohydrochloride (ARG or arginine in the examples)
- 15 • L-Lysine monohydrochloride (LYS or lysine in the examples)
- Trehalose (TRE)
- L-Glycine (GLY or glycine in the examples)
- Saccharose (SAC)
- Sorbitol (SOR)

The following four bacteriostatic agents were evaluated for the multidose formulations:

- Benzyl alcohol (BA)
- m-Cresol (mCr)
- 5 • Phenol (Phe)
- Combination of benzyl alcohol and benzalkonium chloride (BACL)

Example 1 – Investigation on stabilizing agents for monodose formulations

10

Six stabilizing agents (saccharose, arginine, glycine, lysine, sorbitol and trehalose) were tested to deliver stable monodose formulations. The tested formulations are summarized in Table 1.

- 15 Table 1. Composition of several r-hLH monodose formulations containing different stabilizing agents

Components Amount/ml	SAC/250	ARG/250	GLY/250	LYS/250	SOR/250	TRE/250
r-hLH bulk	6 µg	6 µg	6 µg	6 µg	6 µg	6 µg
Na ₂ HPO ₄ 2H ₂ O	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg
NaH ₂ PO ₄ H ₂ O	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg
Methionine	250 µg	250 µg	250 µg	250 µg	250 µg	250 µg
Tween 20	50 µg	50 µg	50 µg	50 µg	50 µg	50 µg
Saccharose	105 mg	-	-	-	-	-
Arginine	-	33 mg	-	-	-	-
Glycine	-	-	23 mg	-	-	-
Lysine	-	-	-	28 mg	-	-
Sorbitol	-	-	-	-	56,5 mg	-
Trehalose	-	-	-	-	-	106 mg
WFI	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml

q.s. = quantity sufficient

- 20 About 40 ml of each solution were prepared, filtered through a 0.22 µm membrane in a 22 ml stainless steel holder, and stored at 2-8°C, +25°C and +40°C in 15 ml plastic tubes. The solutions were tested for protein content (by SE-HPLC), oxidized forms (by RP-HPLC), aggregates (by SE-HPLC) and subunits formation (qualitatively, by SE-HPLC) up to 1 month. The results of the complete panel of tests applied to this
- 25 set of formulations are reported in Tables 2 to 5.

Table 2. Purity by SE-HPLC (storage temperature: +40°C)

Formulation	% heterodimers			% free-subunits			% aggregates		
	T=0	T=1d	T=4d	T=0	T=1d	T=4d	T=0	T=1d	T=4d
SAC/250	96,37	62,27	50,61	0,00	33,71	45,04	3,63	4,02	4,36
GLY/250	95,29	56,55	42,55	0,00	36,42	54,04	4,71	6,43	3,42
ARG/250	95,82	52,91	23,80	0,00	39,83	71,53	4,18	7,27	4,68
LYS/250	95,66	63,26	36,55	0,00	32,57	59,85	4,35	4,18	3,60
SOR/250	95,55	65,87	41,57	0,00	29,64	53,95	4,45	4,49	4,49
TRE/250	92,73	58,22	42,41	0,00	32,62	48,63	7,28	9,16	8,97

T = time; d = day(s)

5 Table 3. Purity by SE-HPLC (storage temperature: +25°C)

Formulation	% heterodimers			% free-subunits			% aggregates		
	T=0	T=7d	T=1M	T=0	T=7d	T=1M	T=0	T=7d	T=1M
SAC/250	96,37	86,52	81,99	0,00	—	12,34	3,63	13,48	5,68
GLY/250	95,29	97,41	78,52	0,00	—	18,64	4,71	2,59	2,84
ARG/250	95,82	98,01	80,80	0,00	—	16,15	4,18	2,00	3,06
LYS/250	95,66	99,20	80,83	0,00	—	17,19	4,35	0,81	1,99
SOR/250	95,55	80,63	26,45	0,00	18,72	72,27	4,45	0,66	1,28
TRE/250	92,73	95,29	74,86	0,00	—	17,24	7,28	4,71	7,91

M = month

Table 4. % oxidised forms by RP-HPLC

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Formulation	12 days (+40°C)	1 Month (+25°C)
SAC/250	7,62	21,84
GLY/250	7,46	n.a.
ARG/250	6,26	9,59
LYS/250	6,68	4,90
SOR/250	6,62	24,58
TRE/250	19,95	10,45

n.a. = not available

Table 5. r-hLH content by SE-HPLC

Formulation	T=0	T=1d (+40°C)	T=4d (+40°C)	T=0	T=1M (+25°C)
SAC/250	5,50	4,20	4,50	5,50	5,40
GLY/250	6,10	5,10	5,10	6,10	5,03
ARG/250	6,30	6,70	6,40	6,30	6,33
LYS/250	6,00	4,80	5,50	6,00	6,02
SOR/250	5,30	4,80	5,10	5,30	4,93
TRE/250	5,40	4,80	5,40	5,40	5,23

15

Based on these results, saccharose, lysine and arginine were selected as best excipients to be further investigated for their compatibility with an increasing amount (250 µg/ml and 500 µg/ml) of methionine as antioxidant. The composition of the tested solutions is reported in Table 6. It should be noted that these solutions have been prepared *ex novo*.

20

Table 6. Composition of r-hLH monodose formulations containing the selected stabilizing agents and different amount of methionine

Components Amount/ml	SAC/250	ARG/250	LYS/250	SAC/500	ARG/500	LYS/500
r-hLH bulk	6 µg	6 µg	6 µg	6 µg	6 µg	6 µg
Na ₂ HPO ₄ 2H ₂ O	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg
NaH ₂ PO ₄ H ₂ O	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg
Methionine	250 µg	250 µg	250 µg	500 µg	500 µg	500 µg
Tween 20	50 µg	50 µg	50 µg	50 µg	50 µg	50 µg
Saccharose	105 mg	-	-	105 mg	-	-
Arginine	-	33 mg	-	-	33 mg	-
Lysine	-	-	28 mg	-	-	28 mg
WFI	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml

- 5 About 100 ml of each solution were prepared, filtered through a 0.22 µm membrane in a 22 ml stainless steel holder, and filled in 1 ml glass syringes to test also the compatibility between the drug substance and the final container (1 ml glass syringe+ plunger). The syringes were stored at 2-8°C, +25°C, +33°C and +40°C. The +33°C storage temperature was introduced in order to have a compromise between a too
- 10 fast (+40°C) and a too low (+25°C) degradation kinetic. The solutions were tested for r-hLH content (by SE-HPLC), oxidized forms (by RP-HPLC), aggregates (by SE-HPLC) and subunits formation (qualitatively, by SE-HPLC) up to 2-3 months.
- The results of the complete panel of test applied to the formulations described in Table 6 are reported in Tables 7 to 11.

15

Table 7. Purity by SE-HPLC (storage temperature: +40°C)

Formulation	% heterodimers		% free-subunits		% aggregates		% heterodimers (2-8°C)	
	T=0	T=4d	T=0	T=4d	T=0	T=4d	T=0	T=13w
SAC/250	99,12	38,98	—	60,29	0,89	0,74	99,12	97,96
ARG/250	98,30	9,16	—	90,03	1,70	0,83	98,30	99,23
LYS/250	98,59	18,27	—	81,02	1,41	0,72	98,59	99,69
SAC/500	99,34	33,96	—	64,11	0,66	1,93	99,34	98,66
ARG/500	98,41	13,49	—	85,94	1,59	0,57	98,41	99,36
LYS/500	98,72	29,07	—	69,89	1,22	1,05	98,72	99,68

w = weeks

20 Table 8. Purity by SE-HPLC (storage temperature: +33°C)

Formulation	% heterodimers					% free-subunits				
	T=0	T=6d	T=4w	T=6w	T=8w	T=0	T=6d	T=4w	T=6w	T=8w
SAC/250	99,12	69,75	65,46	64,82	n.a.	—	29,25	46,04	32,49	n.a.
ARG/250	98,30	65,03	39,69	37,00	31,37	—	32,20	58,05	61,22	63,90
LYS/250	98,52	74,96	53,95	46,28	43,23	—	23,59	44,42	52,58	55,09

Formulation	% heterodimers					% free-subunits				
	T=0	T=6d	T=4w	T=6w	T=8w	T=0	T=6d	T=4w	T=6w	T=8w
SAC/500	99,34	71,26	43,14	27,77	n.a.	—	27,12	50,74	52,34	n.a.
ARG/500	98,41	67,04	38,04	32,62	25,34	—	31,14	58,48	65,37	70,81
LYS/500	98,72	74,03	52,81	49,17	43,88	—	24,88	45,07	49,04	53,24
Formulation	% aggregates									
	T=0		T=6d		T=4w		T=6w		T=8w	
SAC/250	0,89		1,01		n.a.		2,70		n.a.	
ARG/250	1,70		2,77		2,26		1,79		4,74	
LYS/250	1,41		1,45		1,64		1,14		1,69	
SAC/500	0,66		1,63		19,90		6,12		n.a.	
ARG/500	1,59		1,83		3,48		2,02		3,86	
LYS/500	1,22		1,09		2,12		1,79		2,85	

Table 9. Purity by SE-HPLC (storage temperature: +25°C)

Formulation	% heterodimers			% free-subunits			% aggregates		
	T=0	T=4w	T=8w	T=0	T=4w	T=8w	T=0	T=4w	T=8w
SAC/250	99,12	98,36	76,81	—	—	21,05	0,89	1,65	2,15
ARG/250	98,30	79,80	73,39	—	17,89	23,99	1,70	2,32	2,63
LYS/250	98,59	84,15	77,60	—	13,78	21,23	1,41	n.a.	1,18
SAC/500	99,34	n.a.	61,24	—	n.a.	33,50	0,66	n.a.	5,26
ARG/500	98,41	n.a.	69,45	—	n.a.	27,93	1,59	n.a.	2,63
LYS/500	98,72	n.a.	78,56	—	n.a.	19,81	1,22	n.a.	1,64

5 Table 10. % oxidised forms by RP-HPLC

Formulation	4 days (+40°C)	8 weeks (+25°C)	4 weeks (+33°C)	6 weeks (+33°C)
SAC/250	6,13	7,32	9,72	20,13
SAC/500	17,37	35,78	39,12	45,93
ARG/250	6,29	5,96	9,72	7,02
ARG/500	4,69	5,67	6,69	8,10
LYS/250	5,24	5,05	8,00	8,63
LYS/500	3,25	4,02	5,71	6,84

Table 11. r-hLH content by SE-HPLC

Formulation	T=0	T=4days (+40°C)	T=8weeks (+25°C)	T=4weeks (+33°C)	T=6weeks (+33°C)	T=8weeks (+33°C)
SAC/250	5,03	3,09	5,09	5,31	4,74	n.a.
SAC/500	6,27	4,92	6,00	5,96	5,63	6,20
ARG/250	5,90	3,74	5,50	5,63	5,34	5,17
ARG/500	5,00	3,36	4,25	5,05	n.a.	n.a.
LYS/250	6,14	4,54	5,74	5,83	5,58	5,84
LYS/500	6,08	3,91	5,61	5,73	5,49	5,48

10

Based on these results saccharose showed a higher compatibility with 250 µg/ml of methionine, lysine with 500 µg/ml of methionine and arginine with both concentrations of methionine.

Example 2 – Compatibility of r-hLH monodose liquid formulations with FSH multidose liquid formulation

In view of the results disclosed in Example 1, the formulations described in Table 6 were mixed with an FSH formulation (i.e. Gonal-F® multidose liquid formulation) and tested after 24 h of contact at 25°C according to the methods below:

- RP-HPLC for r-hFSH and r-hLH α -subunit oxidised forms (in Table 12 under % purity the non-oxidised forms are reported)
- SE-HPLC for aggregates quantitation
- RP-HPLC for r-hFSH and r-hLH titre,
- r-hFSH and r-hLH *in vivo* bioassay
- SDS-PAGE for r-hFSH and r-hLH free-subunits and aggregates quantitation (data not reported),
- Visual appearance.

The results are reported in Tables 12 to 14.

Table 12. Purity by RP-HPLC and % aggregates by SE-HPLC

Formulation	% purity		% aggregates	
	T=0	T=24 h	T=0	T=24 h
SAC/250	98,53	98,61	0,00	0,00
ARG/250	97,29	99,17	0,00	0,72
LYS/250	97,81	97,73	0,00	0,37
SAC/500	98,50	98,17	0,00	0,39
ARG/500	97,75	97,47	0,00	0,69
LYS/500	97,04	96,74	0,00	0,33

h = hours

Table 13. FSH and LH content

Formulation	r-hFSH content*		r-hLH content [#]	
	T=0	T=24 h	T=0	T=24 h
SAC/250	16,14	16,14	3,67	3,61
ARG/250	15,45	16,01	4,15	4,21
LYS/250	15,35	15,68	4,07	4,15
SAC/500	16,29	16,91	3,71	3,70
ARG/500	16,03	15,80	4,15	4,05
LYS/500	15,46	15,69	4,05	4,17

* FSH theoretical: 200 IU=15,48 µg/ml; [#] LH theoretical: 100 IU=4 µg/ml

Table 14. Bioassay

Formulation	r-hFSH activity		r-hLH activity	
	T=0	T=24 h	T=0	T=24 h
SAC/250	174,60	183,70	108,00	109,00

All the tested monodose formulations were compatible with the Gonal-F® multidose formulation as there was:

- No loss in FSH and LH content,
- No oxidation,
- 5 - No aggregation and free-subunits dissociation (by SDS-PAGE)
- No loss in LH and FSH bioactivity.

Example 3 – Stability study of monodose liquid formulations

10 On the basis of the results of the previous examples, the formulations described in Table 6, except those containing arginine, were prepared at two different r-hLH concentrations (6 µg/ml and 12 µg/ml), stored at 2-8°C and +25°C and tested according to a rigorous stability plan and according to the following analytical methods:

- 15 - RP-HPLC for LH content
- RP-HPLC for α-subunit oxidised forms
- SDS-PAGE for free-subunits and aggregates
- Bioassay
- pH of solution
- 20 - Visual appearance

A statistical analysis was performed on the results of all the parameters that were identified as stability indicating with the aid of Stabileo 1.1 software.

25 *r-hLH concentration (by RP-HPLC) upon storage at 2-8°C and 25±2°C*

No statistically significant loss in protein concentration was observed for the formulations after 6-months storage at 2-8°C whatever the r-hLH concentration (6 µg/ml and 12 µg/ml). A common decrease of 0.4 µg/month was observed after 6-
30 months storage at 25±2°C.

r-hLH bioactivity (bioassay) upon storage at 2-8°C and 25±2°C

No relevant loss in bioactivity was observed for the formulations after 6-months
35 storage at 2-8°C and at 25±2°C whatever the r-hLH concentration (6 µg/ml and 12 µg/ml).

% subunits and aggregates by SDS-PAGE

The percentage of aggregates by SDS-PAGE remains below 2% for both concentrations after 6-months storage at 2-8°C and 25°C. The percentage of subunits by SDS-PAGE remains below 2% for both concentrations after 6-months storage at 2-8°C. A common decrease of about 5%/month was observed after 6-months storage at 25°C±2°C.

% oxidised forms by RP-HPLC

An increase of about 0.4%/month was measured after 6-months storage at 2-8°C and an increase in the range 0.6-1.4%/month at 25°C±2°C.

pH and appearance

No change in the appearance (colour, clarity, visible particles) nor in the pH was observed during manufacturing and upon storage.

Example 4 – Compatibility of stabilizing and bacteriostatic agents for multidose formulations

Six stabilizing agents (saccharose, arginine, glycine, lysine, sorbitol and trehalose) were tested for compatibility with bacteriostatic agents to deliver stable multidose formulations. The tested formulations are summarized in Table 15.

Table 15. Composition of several r-hLH multidose formulations containing different combinations of stabilizing and bacteriostatic agents

Components Amount/ml	SAC/250 BA	SAC/250 mCr	SAC/250 Phe	ARG/250 BA	ARG/250 mCr	ARG/250 Phe
r-hLH bulk	25 µg	25 µg	25 µg	25 µg	25 µg	25 µg
Na ₂ HPO ₄ 2H ₂ O	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg
NaH ₂ PO ₄ H ₂ O	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg
Methionine	250 µg	250 µg	250 µg	250 µg	250 µg	250 µg
Tween 20	50 µg	50 µg	50 µg	50 µg	50 µg	50 µg
Saccharose	105 mg	105 mg	105 mg	-	-	-
Arginine	-	-	-	33 mg	33 mg	33 mg
Benzyl Alcohol	0.90%	-	-	0.90%	-	-
m-Cresol	-	0.30%	-	-	0.30%	-
Phenol	-	-	0.50%	-	-	0.50%
WFI	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml

Components Amount/ml	LYS/250 BA	LYS/250m Cr	LYS/250 Phe	SOR/250 BA	SOR/250 mCr	SOR/250 Phe
r-hLH bulk	25 µg	25 µg	25 µg	25 µg	25 µg	25 µg
Na ₂ HPO ₄ 2H ₂ O	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg
NaH ₂ PO ₄ H ₂ O	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg
Methionine	250 µg	250 µg	250 µg	250 µg	250 µg	250 µg
Tween 20	50 µg	50 µg	50 µg	50 µg	50 µg	50 µg
Lysine	28 mg	28 mg	28 mg	-	-	-
Sorbitol	-	-	-	56.5 mg	56.5 mg	56.5 mg
Benzyl Alcohol	0.90%	-	-	0.90%	-	-
m-Cresol	-	0.30%	-	-	0.30%	-
Phenol	-	-	0.50%	-	-	0.50%
WFI	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml
Components Amount/ml	GLY/250/ BA	GLY/250/ mCr	GLY/250/ Phe	TRE/250/ BA	TRE/250/ mCr	TRE/250/ Phe
r-hLH bulk	25 µg	25 µg	25 µg	25 µg	25 µg	25 µg
Na ₂ HPO ₄ 2H ₂ O	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg
NaH ₂ PO ₄ H ₂ O	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg
Methionine	250 µg	250 µg	250 µg	250 µg	250 µg	250 µg
Tween 20	50 µg	50 µg	50 µg	50 µg	50 µg	50 µg
Glycine	23 mg	23 mg	23 mg	-	-	-
Trehalose	-	-	-	106 mg	106 mg	106 mg
Benzyl Alcohol	0.90%	-	-	0.90%	-	-
m-Cresol	-	0.30%	-	-	0.30%	-
Phenol	-	-	0.50%	-	-	0.50%
WFI	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml

About 40 ml of each solution were prepared, filtered through a 0.22 µm membrane and stored at 2-8°C, +25°C and +40°C in 15 ml plastic tubes. The solutions were tested for protein content (by Size Exclusion-HPLC, SE-HPLC, data not shown),

- 5 oxidized forms (by Reverse Phase-HPLC, RH-HPLC), aggregates (by SE-HPLC) and subunits formation (qualitatively, by SE-HPLC) up to 1 month. The results of the complete panel of tests applied to this set of formulations are reported in Tables 16 to 18. All the solutions containing m-Cresol (mCr) became opalescent even immediately after manufacturing, due to an incompatibility between the bacteriostatic agent and
- 10 the surfactant (Tween 20).

Table 16. Purity by SE-HPLC (storage temperature: +40°C)

Formulation	% heterodimers			% free-subunits			% aggregates		
	T=0	T=1d	T=4d	T=0	T=1d	T=4d	T=0	T=1d	T=4d
SAC/250/mCr	98,91	89,61	52,93	—	n.a.	23,67	1,10	10,38	23,41
SAC/250/BA	95,50	74,04	51,52	—	22,65	45,12	4,50	3,33	3,36
SAC/250/Phe	98,34	74,20	67,13	—	19,63	23,71	1,66	5,57	9,17
GLY/250/mCr	97,95	96,54	86,76	—	9,55	n.a.	2,05	3,14	n.a.
GLY/250/BA	96,08	59,58	41,59	—	37,82	56,45	3,92	2,61	1,97
GLY/250/Phe	96,49	74,71	86,06	—	20,01	9,22	2,51	5,28	4,73
ARG/250/mCr	97,94	70,34	44,39	—	28,39	53,33	2,06	1,29	2,29
ARG/250/BA	91,44	35,99	15,91	—	58,99	79,65	8,57	5,03	4,45
ARG/250/Phe	95,31	63,27	29,12	—	33,75	67,18	4,69	2,98	3,70

Formulation	% heterodimers			% free-subunits			% aggregates		
	T=0	T=1d	T=4d	T=0	T=1d	T=4d	T=0	T=1d	T=4d
LYS/250/mCr	98,19	81,57	65,68	—	17,99	33,10	1,81	n.a.	1,22
LYS/250/BA	94,75	47,94	33,61	—	48,59	63,14	5,26	3,48	3,26
LYS/250/Phe	97,91	72,68	56,73	—	26,18	42,26	2,09	1,16	1,01
SOR/250/mCr	90,46	86,33	58,78	—	n.a.	18,87	9,55	13,68	22,36
SOR/250/BA	94,08	62,71	45,27	—	64,17	47,22	5,93	3,13	7,52
SOR/250/Phe	94,74	70,58	55,75	—	22,30	32,76	5,26	7,13	11,50
TRE/250/mCr	86,07	77,26	40,60	—	n.a.	20,87	13,93	22,74	38,53
TRE/250/BA	94,74	69,53	42,19	—	21,36	44,22	5,26	9,12	12,88
TRE/250/Phe	91,05	85,29	54,89	—	n.a.	24,28	8,95	14,72	20,83

Table 17. Purity by SE-HPLC (storage temperature: +25°C)

Formulation	% heterodimers			% free-subunits			% aggregates		
	T=0	T=7d	T=1M	T=0	T=7d	T=1M	T=0	T=7d	T=1M
SAC/250/mCr	98,91	53,76	69,30	—	31,19	15,13	1,10	14,34	15,57
SAC/250/BA	95,50	95,04	75,84	—	—	14,66	4,50	4,96	9,50
SAC/250/Phe	98,34	93,82	85,49	—	—	—	1,66	6,18	14,51
GLY/250/mCr	97,95	97,00	95,39	—	—	—	2,05	3,00	4,62
GLY/250/BA	96,08	97,63	68,21	—	—	28,42	3,92	2,37	2,17
GLY/250/Phe	96,49	97,44	93,19	—	—	—	2,51	2,57	6,10
ARG/250/mCr	97,94	98,73	78,33	—	—	19,73	2,06	1,28	1,94
ARG/250/BA	91,44	97,38	79,79	—	—	18,91	8,57	2,63	1,31
ARG/250/Phe	95,31	98,33	82,18	—	—	16,59	4,69	1,67	1,23
LYS/250/mCr	98,19	98,28	95,38	—	—	—	1,81	1,73	4,27
LYS/250/BA	94,75	97,39	82,54	—	—	14,66	5,26	2,61	2,80
LYS/250/Phe	97,91	99,55	98,61	—	—	—	2,09	0,46	1,40
SOR/250/mCr	90,46	82,65	64,24	—	—	—	9,55	17,23	35,76
SOR/250/BA	94,08	95,93	68,59	—	—	21,99	5,93	4,08	9,43
SOR/250/Phe	94,74	92,55	81,96	—	—	—	5,26	7,46	18,05
TRE/250/mCr	86,07	75,78	45,61	—	—	—	13,93	24,23	54,40
TRE/250/BA	94,74	91,22	67,28	—	—	13,75	5,26	8,79	18,97
TRE/250/Phe	91,05	84,53	66,79	—	—	—	8,95	15,47	33,22

Table 18. % oxidised forms by RP-HPLC

Formulation	12 days (+40°C)	1 Month (+25°C)
SAC/250/mCr	51,14	16,62
SAC/250/BA	14,07	8,83
SAC/250/Phe	25,55	17,54
GLY/250/mCr	59,29	5,43
GLY/250/BA	10,53	4,44
GLY/250/Phe	34,86	16,06
ARG/250/mCr	65,27	3,13
ARG/250/BA	7,12	5,65
ARG/250/Phe	15,49	11,27
LYS/250/mCr	44,05	16,36
LYS/250/BA	13,69	9,30
LYS/250/Phe	56,55	17,86
SOR/250/mCr	58,29	25,68
SOR/250/BA	13,51	8,49
SOR/250/Phe	30,78	17,75
TRE/250/mCr	45,84	26,36
TRE/250/BA	19,12	11,41
TRE/250/Phe	23,87	27,76

Based on these results, saccharose, lysine and arginine were selected as the best stabilizing agents to be further investigated for their compatibility with an increased amount of methionine (500 µg/ml). Benzyl alcohol, phenol and a combination of benzyl alcohol and benzalkonium chloride were selected as bacteriostatic agents.

- 5 The tested formulations are summarized in Table 19.

Table 19. Composition of several r-hLH multidose formulations containing the selected stabilizing agents and different bacteriostatic agents

Components Amount/ml	SAC/500 BA	SAC/500 BACL	SAC/500 Phe	ARG/500 BA	ARG/500 BACL	ARG/500 Phe
r-hLH bulk	25 µg	25 µg	25 µg	25 µg	25 µg	25 µg
Na ₂ HPO ₄ 2H ₂ O	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg	1.65 mg
NaH ₂ PO ₄ H ₂ O	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg	0.104 mg
Methionine	500 µg	500 µg	500 µg	500 µg	500 µg	500 µg
Tween 20	50 µg	50 µg	50 µg	50 µg	50 µg	50 µg
Saccharose	105 mg	105 mg	105 mg	-	-	-
Arginine	-	-	-	33 mg	33 mg	33 mg
Benzyl Alcohol	1.2%	0.30%	-	1.2%	0.30%	-
Benzalkonium chloride	-	0.001%	-	-	0.001%	-
Phenol	-	-	0.50%	-	-	0.50%
WFI	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml	q.s. to 1 ml
Components Amount/ml	LYS/500/BA		LYS/500/BACL		LYS/500/Phe	
r-hLH bulk	25 µg		25 µg		25 µg	
Na ₂ HPO ₄ 2H ₂ O	1.65 mg		1.65 mg		1.65 mg	
NaH ₂ PO ₄ H ₂ O	0.104 mg		0.104 mg		0.104 mg	
Methionine	500 µg		500 µg		500 µg	
Tween 20	50 µg		50 µg		50 µg	
Lysine	28 mg		28 mg		28 mg	
Benzyl Alcohol	1.2%		0.30%		-	
Benzalkonium chloride	-		0.001%		-	
Phenol	-		-		0.50%	
WFI	q.s. to 1 ml		q.s. to 1 ml		q.s. to 1 ml	

- 10 About 100 ml of each solution were prepared, filtered through a 0.22 µm membrane and filled in 3 ml cartridges. The following primary packaging was used:
- 3 ml glass cartridges (Nuova Ompi) siliconised
 - Crimp caps code CAP J 3ML L1H075-1-H1B FM 257/2 (Helvoet Pharma)
 - Coated plungers: Helvoet V9282 FM257/2 Omniflex coated.
- 15 The cartridges were stored at 2-8°C, +25°C and +40°C to be tested for protein content (by SE-HPLC), oxidized forms (by RH-HPLC), aggregates (by SE-HPLC) and subunits formation (qualitatively, by SE-HPLC) up to 2 months.

The results of the complete panel of tests applied to this second set of formulations are reported in Tables 20 to 26.

Table 20. Purity by SE-HPLC (storage temperature: +40°C)

Formulation	% heterodimers		% free-subunits		% aggregates	
	T=0	T=3 days	T=0	T=3 days	T=0	T=3 days
SAC/500/BA	100,00	60,01	—	39,55	—	0,44
SAC/500/BACL	100,00	67,39	—	32,34	—	0,27
SAC/500/Phe	99,66	65,01	—	31,27	0,34	3,72
ARG/500/BA	99,91	45,63	—	54,65	0,10	—
ARG/500/BACL	99,93	44,67	—	55,33	0,07	—
ARG/500/Phe	99,93	61,97	—	38,03	0,08	—
LYS/500/BA	99,93	61,71	—	38,30	0,08	—
LYS/500/BACL	99,95	61,33	—	38,68	0,06	—
LYS/500/Phe	99,93	76,03	—	23,59	0,08	0,39

Table 21. Purity by SE-HPLC (storage temperature: +33°C)

Formulation	% heterodimers				% free-subunits			% aggregates	
	T=0	T=3w	T=4w	T=8w	T=0	T=3w	T=4w	T=0	T=3w
SAC/500/BA	100,00	76,56	71,22	59,22	—	22,03	26,35	/	1,43
SAC/500/BACL	100,00	81,49	78,95	63,53	—	17,26	20,10	/	1,26
SAC/500/Phe	99,66	76,24	67,13	51,10	—	14,26	18,03	0,34	9,51
ARG/500/BA	99,91	71,90	71,52	61,41	—	27,89	27,73	0,10	0,21
ARG/500/BACL	99,93	69,08	66,86	57,22	—	30,67	32,94	0,07	0,26
ARG/500/Phe	99,93	86,11	76,74	75,51	—	13,76	22,55	0,08	0,14
LYS/500/BA	99,93	83,09	76,72	67,30	—	16,76	22,50	0,08	0,16
LYS/500/BACL	99,95	81,03	75,21	70,20	—	18,88	24,64	0,06	0,09
LYS/500/Phe	99,93	87,29	80,51	68,85	—	12,10	18,53	0,08	0,62

5

Table 22. Purity by SE-HPLC (storage temperature: +25°C)

Formulation	% heterodimers			% free-subunits		% aggregates
	T=0	T=4w	T=8w	T=0	T=4w	T=0
SAC/500/BA	100,00	83,85	82,87	—	14,69	—
SAC/500/BACL	100,00	86,30	85,78	—	13,06	—
SAC/500/Phe	99,66	79,35	75,26	—	13,39	0,34
ARG/500/BA	99,91	82,32	82,13	—	17,39	0,10
ARG/500/BACL	99,93	80,42	80,77	—	19,25	0,07
ARG/500/Phe	99,93	84,59	80,95	—	15,13	0,08
LYS/500/BA	99,93	85,04	85,10	—	13,78	0,08
LYS/500/BACL	99,95	84,24	86,22	—	15,42	0,06
LYS/500/Phe	99,93	85,41	81,22	—	13,68	0,08

Table 23. % oxidised forms by RP-HPLC (storage temperature: +33°C)

Formulation	T=0	T=3 weeks	T=8 weeks	T=13 weeks
SAC/500/BA	1,12	5,21	4,21	4,26
SAC/500/BACL	0,90	5,59	4,90	4,46
SAC/500/Phe	0,99	8,28	6,01	16,14
ARG/500/BA	0,69	4,52	4,13	2,65
ARG/500/BACL	1,13	4,57	3,57	2,97
ARG/500/Phe	0,94	4,60	5,24	4,82
LYS/500/BA	1,53	5,71	16,34	8,77
LYS/500/BACL	1,60	4,49	5,83	5,70
LYS/500/Phe	0,66	9,66	34,97	25,51

Table 24. % oxidised forms by RP-HPLC (storage temperature: +25°C)

Formulation	T=0	T=4 weeks	T=8 weeks	T=13 weeks
SAC/500/BA	1,12	2,04	2,19	2,44
SAC/500/BACL	0,90	2,10	2,31	2,23
SAC/500/Phe	0,99	2,68	5,72	7,50
ARG/500/BA	0,69	1,40	1,61	1,65
ARG/500/BACL	1,13	1,46	2,31	1,89
ARG/500/Phe	0,94	1,31	2,77	2,35
LYS/500/BA	1,53	2,51	3,35	4,73
LYS/500/BACL	1,60	1,96	3,14	3,47
LYS/500/Phe	0,66	4,05	11,38	22,52

Table 25. % oxidised forms by RP-HPLC

Formulation	T=0	T=13 weeks (+5°C)	T=0	T=3 days (+40°C)
SAC/500/BA	1,12	1,11	1,12	1,16
SAC/500/BACL	0,90	1,29	0,90	1,25
SAC/500/Phe	0,99	2,51	0,99	1,47
ARG/500/BA	0,69	1,03	0,69	0,64
ARG/500/BACL	1,13	1,16	1,13	1,06
ARG/500/Phe	0,94	2,02	0,94	1,47
LYS/500/BA	1,53	2,03	1,53	1,84
LYS/500/BACL	1,60	1,56	1,60	1,58
LYS/500/Phe	0,66	n.a.	0,66	3,10

5 Table 26. r-hLH content by SE-HPLC

Formulation	T=0	T=3d (+40°C)	T=3w (+33°C)	T=4w (+33°C)	T=8w (+33°C)	T=4w (+25°C)	T=8w (+25°C)
SAC/500/BA	20,38	21,70	21,05	22,54	16,47	23,70	18,43
SAC/500/BACL	21,73	22,06	21,31	22,64	18,77	23,65	18,10
SAC/500/Phe	21,24	21,49	21,02	22,53	20,71	22,66	17,23
ARG/500/BA	22,23	18,86	19,52	24,64	16,14	25,26	17,69
ARG/500/BACL	22,38	21,48	21,71	23,52	16,86	24,18	19,34
ARG/500/Phe	21,34	22,38	20,60	23,09	16,99	24,41	16,32
LYS/500/BA	20,33	20,93	18,85	22,10	14,93	24,24	17,36
LYS/500/BACL	23,83	21,78	20,68	23,44	15,50	24,07	18,60
LYS/500/Phe	21,34	21,05	20,20	21,89	16,50	22,99	16,56

These results confirmed the compatibility of the stabilizing agents with an increased amount of methionine (see Example 1). In addition the results showed that the bacteriostatic agents are compatible with 500 µg/ml of methionine.

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Example 5 – Compatibility of r-hLH multidose liquid formulations with FSH multidose liquid formulation

Based on the results of Example 4, formulations of r-hLH in 3 ml cartridges were mixed with a FSH formulation (i.e. Gonal-F® multidose liquid formulation) and tested after 24 h of contact at 25°C according to the methods below:

15

- SE-HPLC for purity

- RP-HPLC for r-hFSH and r-hLH titre,
- RP-HPLC for r-hFSH and r-hLH α -subunit oxidised forms,
- r-hFSH and r-hLH *in vivo* bioassay
- SDS-PAGE for r-hFSH and r-hLH free-subunits and aggregates quantitation (data not reported),
- pH of solution,
- Visual appearance.

The results are reported in Tables 27 to 29.

10 Table 27. Purity by SE-HPLC

Formulation	% heterodimers		% aggregates	
	T=0	T=24 h	T=0	T=24 h
SAC/500/BA	98,28	98,41	0,30	1,09
SAC/500/BACL	98,85	99,66	0,00	0,43
SAC/500/Phe	98,18	97,68	0,27	0,54
ARG/500/BA	98,31	98,61	0,28	0,98
ARG/500/BACL	97,68	98,52	0,33	0,53
ARG/500/Phe	97,89	97,73	0,44	0,77
LYS/500/BA	97,35	97,51	0,38	1,09
LYS/500/BACL	97,97	98,82	0,28	0,81
LYS/500/Phe	98,29	98,40	0,37	1,01

Table 28. FSH and LH content

Formulation	r-hFSH content*		r-hLH content [#]	
	T=0	T=24 h	T=0	T=24 h
SAC/500/BACL	29,50	31,15	7,20	8,31
SAC/500/Phe	30,88	31,28	7,34	7,54
ARG/500/BA	30,46	31,00	8,02	8,44
ARG/500/BACL	32,23	32,84	7,12	7,40
ARG/500/Phe	30,07	31,06	7,78	8,05
LYS/500/BA	30,91	32,11	6,88	7,25
LYS/500/BACL	28,57	31,66	6,64	7,62
LYS/500/Phe	28,75	31,25	6,80	7,47

* FSH theoretical: 200 IU=30,96 μ g/ml; [#] LH theoretical: 100 IU=8 μ g/ml

15 Table 29. Bioassay

Formulation	r-hFSH activity		r-hLH activity	
	T=0	T=24 h	T=0	T=24 h
SAC/500/BACL	365,7	355,0	227,0	212,7
ARG/500/Phe	387,5	393,0	231,4	243,0

All the tested multidose formulations are compatible with the Gonal-F[®] multidose formulation as there was:

- No loss in FSH and LH content,
- No oxidation,
- No aggregation (by SE-HPLC)

- No aggregates and subunits formation by SDS-PAGE
- No loss in LH and FSH bioactivity.

Example 6 – Compatibility with primary packaging

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In order to test the compatibility between the r-hLH formulations and the final containers (syringes and cartridges), a matrix study was put in place to test the following parameters:

- Effect of r-hLH concentration: 6 µg/ml, 12 µg/ml, 24 µg/ml,
- 10 - Effect of filling volume: 0.25 ml, 0.5 ml, 1 ml,
- Type of container: 1ml glass syringe and 3 ml glass cartridge,
- Effect of bacteriostatic agent: formulations prepared with and without the bacteriostatic agents and stored in 3 ml cartridges.

One formulation (SAC/500/BACL) was prepared at the different r-hLH strengths and filled in the containers at different filling volumes. The batches were compared qualitatively by SE-HPLC up to 1 week at +33°C. The results are reported in Figure 1 (A-F). In Figure 2 (A-B) the percentage of heterodimers by SE-HPLC on the formulation with and without the combination of bacteriostatic agents are compared.

Looking at the results in Figures 1 and 2, the following conclusions can be drawn:

- 20 - No impact of container on formulations at 12 µg/ml and 24 µg/ml
- No impact of filling volume on formulations at 12 µg/ml and 24 µg/ml
- Positive effect of increased concentration on subunits dissociation
- No difference between batches with or without bacteriostatics
- Higher stability is related to higher r-hLH concentration.

25

Example 7 – Stability study of multidose liquid formulations

On the basis of the results of the previous examples, four formulations described in Table 19 (SAC/500/BACL, LYS/500/BA, LYS/500/BACL and ARG/500/Phe) were stored at 2-8°C and +25°C and tested according to a rigorous stability plan and according to the following analytical methods:

- RP-HPLC for LH content
- RP-HPLC for α-subunit oxidised forms
- RP-HPLC for phenol content
- 35 - RP-HPLC for benzalkonium chloride content
- RP-HPLC for benzyl alcohol content
- SDS-PAGE for free-subunits and aggregates

- Bioassay
- pH of solution
- Visual appearance

5 The lysine and arginine amount in the above formulations was adjusted (i.e. L-lysine monohydrochloride to 28.5 mg and L-arginine monohydrochloride to 31.5 mg) to optimize the isotonicity of the formulations.

A statistical analysis was performed on the results of all the parameters that were identified as stability indicating with the aid of Stabileo 1.1 software.

10

r-hLH concentration (by RP-HPLC) upon storage at 2-8°C and 25±2°C

No statistically significant loss in protein concentration was observed for the formulations after 12-months storage at 2-8°C. A common decrease of 0.4 µg/month was observed after 6-months storage at 25±2°C.

15

r-hLH bioactivity (bioassay) upon storage at 2-8°C and 25±2°C

No relevant loss in bioactivity was observed for the formulations after 12-months storage at 2-8°C and at 25±2°C.

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% subunits and aggregates by SDS-PAGE

The percentage of aggregates by SDS-PAGE remains below 2% for all formulations after 12-months storage at 2-8°C and 25°C.

25

The percentage of subunits by SDS-PAGE increases above 32% after 6 months at 25°C for all the formulations.

% oxidised forms by RP-HPLC

30

SAC/500/BACL and ARG/500/Phe present the lower increase of oxidised forms.

pH and appearance

35 No change in the appearance (colour, clarity, visible particles) nor in the pH was observed during manufacturing and upon storage.

Bacteriostatic agents content

The target amounts of phenol and benzyl alcohol were detected and no losses were measured from 6-months to 12-months stability checkpoints. An amount of benzalkonium chloride below target (6 µg/ml instead of 10 µg/ml) was measured in the SAC/500/BACL formulation at 6-months stability checkpoint.

Efficacy of bacteriostatic agents

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The results of the bacteriostatic efficacy tests, performed at T=0 and repeated after 9 months on the formulations are reported in Tables 30 to 33.

Table 30. Results of bacteriostatic efficacy test on multidose formulation SAC/500/BACL

Microorganism	Log reduction vs T=0				
	T=6 h	T=24 h	T=7 days	T=14 days	T=28 days
<i>Staphylococcus aureus</i>	>3	>3	>3	>3	no reduction
<i>Pseudomonas aeruginosa</i>	>3	>3	>3	>3	no reduction
<i>Escherichia coli</i>	no tested	n.t.	>3	>3	no reduction
<i>Candida albicans</i>	n.t.	n.t.	>3	>3	no increase
<i>Aspergillus niger</i>	n.t.	n.t.	1.8	>3	no increase

Table 31. Results of bacteriostatic efficacy test on multidose formulation LYS/500/BACL

Microorganism	Log reduction vs T=0				
	T=6 h	T=24 h	T=7 days	T=14 days	T=28 days
<i>Staphylococcus aureus</i>	>3	>3	>3	>3	no reduction
<i>Pseudomonas aeruginosa</i>	>3	>3	>3	>3	no reduction
<i>Escherichia coli</i>	not tested	n.t.	>3	>3	no reduction
<i>Candida albicans</i>	n.t.	n.t.	>3	>3	no increase
<i>Aspergillus niger</i>	n.t.	n.t.	3.3	>3	no increase

Table 32. Results of bacteriostatic efficacy test on multidose formulation LYS/500/BA

Microorganism	Log reduction vs T=0				
	T=6 h	T=24 h	T=7 days	T=14 days	T=28 days
<i>Staphylococcus aureus</i>	1.05	>3	>3	>3	no reduction
<i>Pseudomonas aeruginosa</i>	>3	>3	>3	>3	no reduction
<i>Escherichia coli</i>	not tested	n.t.	>3	>3	no reduction
<i>Candida albicans</i>	n.t.	n.t.	>3	>3	no increase

<i>Aspergillus niger</i>	n.t.	n.t.	>3	>3	no increase
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Table 33. Results of bacteriostatic efficacy test on multidose formulation ARG/500/Phe

Microorganism	Log reduction vs T=0				
	T=6 h	T=24 h	T=7 days	T=14 days	T=28 days
<i>Staphylococcus aureus</i>	0.73	>3	>3	>3	no reduction
<i>Pseudomonas aeruginosa</i>	>3	>3	>3	>3	no reduction
<i>Escherichia coli</i>	not tested	n.t.	>3	>3	no reduction
<i>Candida albicans</i>	n.t.	n.t.	>3	>3	no increase
<i>Aspergillus niger</i>	n.t.	n.t.	>3	>3	no increase

The formulations containing the combination of 0.3% benzyl alcohol + 0.001% benzalkonium chloride, and saccharose or lysine (SAC/500/BACL and LYS/500/BACL) met Criteria A of the European Pharmacopeia even if the amount of benzalkonium chloride was below the target (6 µg/ml instead of 10 µg/ml).

The formulation containing 0.5% phenol (ARG/500/Phe) and the formulation containing 1.2% benzyl alcohol (LYS/500/BA) met Criteria B of the European Pharmacopeia.

Conclusions

All multidose formulations showed a good stability profile after 12 months storage at 2-8°C.

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Claims

1. A liquid formulation containing luteinizing hormone (LH) or a variant thereof characterized in that said formulation comprises a phosphate buffer and a
5 stabilizing amount of arginine or salts thereof at a concentration of 10 to 50 mg/ml and/or lysine or salts thereof at a concentration of 10 to 50 mg/ml.
2. The formulation according to claim 1, wherein the luteinizing hormone (LH) is human luteinizing hormone (hLH).
3. The formulation according to claim 1 or 2, wherein the luteinizing hormone
10 (LH) is recombinant human luteinizing hormone (r-hLH).
4. The formulation according to claim 1 or 2, wherein the luteinizing hormone (LH) is urinary human luteinizing hormone (u-hLH).
5. The formulation according to any of the preceding claims, wherein the luteinizing hormone (LH) is present at a concentration of 1 to 50 µg/ml.
- 15 6. The formulation according to any of the preceding claims, wherein the phosphate buffer is present at a concentration of 1 to 100 mM.
7. The formulation according to any of the preceding claims, wherein the phosphate buffer is preferably sodium phosphate buffer.
8. The formulation according to any of the preceding claims, further comprising
20 a surfactant.
9. The formulation according to claim 8, wherein the surfactant is Polysorbate 20.
10. The formulation according to claim 9, wherein Polysorbate 20 is present at a concentration of 0.01 to 10 mg/ml.
- 25 11. The formulation according to any of the preceding claims, further comprising methionine.
12. The formulation according to claim 11, wherein methionine is present at a concentration of 0.01 to 5.0 mg/ml.

13. The formulation according to any of the preceding claims, further comprising a bacteriostatic agent.
14. The formulation according to claim 13, wherein the bacteriostatic agent is selected from any of benzyl alcohol, phenol and a combination of benzyl alcohol and benzalkonium chloride.
15. The formulation according to claim 13 or 14, wherein the bacteriostatic agent is present at a concentration of 0.005 to 15 mg/ml.
16. The formulation according to any of the preceding claims, further comprising water for injection.
17. The formulation according to any of the preceding claims having a pH in the range of 7.5 to 8.5.
18. The formulation according to any of the preceding claims comprising at least 25 µg/ml of r-hLH, 1.65 mg/ml of Na₂HPO₄ 2H₂O, 0.104 mg/ml of NaH₂PO₄ H₂O, 31.5 mg/ml of L-arginine monohydrochloride, 0.05 mg/ml of Polysorbate 20, 0.5 mg/ml of methionine and 5 mg/ml of phenol.
19. The formulation according to any of claims 1 to 17 comprising at least 25 µg/ml of r-hLH, 1.65 mg/ml of Na₂HPO₄ 2H₂O, 0.104 mg/ml of NaH₂PO₄ H₂O, 28.5 mg/ml of lysine monohydrochloride, 0.05 mg/ml of Polysorbate 20, 0.5 mg/ml of methionine and 12 mg/ml of benzylalcohol.
20. The formulations according to any of claims 1 to 17 comprising at least 25 µg/ml of r-hLH, 1.65 mg/ml of Na₂HPO₄ 2H₂O, 0.104 mg/ml of NaH₂PO₄ H₂O, 28.5 mg/ml of lysine monohydrochloride, 0.05 mg/ml of Polysorbate 20, 0.5 mg/ml of methionine 3 mg/ml of benzylalcohol and 0.01 mg/ml of benzalkonium chloride.
21. A pharmaceutical composition comprising the formulation according to any of claims 1 to 20.
22. A form of presentation of the formulation according to any of claims 1 to 20 hermetically sealed under sterile conditions in a container suitable for storage prior to use.

23. Process for the preparation of the formulation according to any of claims 1 to 20, comprising the dilution of the luteinizing hormone (LH) with a solution of excipients.
- 5 24. Use of the formulation according to any of claims 1 to 20 for the preparation of a medicament.
25. Use of the formulation according to any of claims 1 to 20 for the preparation of a medicament for the treatment of infertility in women and/or men.
- 10 26. Use of the formulation according to any of claims 1 to 20 in combination with a liquid formulation comprising follicle-stimulating hormone (FSH) or a variant thereof.
27. The formulation according to any of claims 1 to 20 for the treatment of infertility in women and/or men.