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(72) Inventeurs/Inventors:
KALLMEYER, GEORG, DE;
WINTER, GERHARD, DE;
KLESSEN, CHRISTIAN, DE;
WOOG, HEINRICH, DE

(73) Propriétaire/Owner:
ROCHE DIAGNOSTICS GMBH, DE

(74) Agent: NORTON ROSE FULBRIGHT CANADA
LLP/S.E.N.C.R.L., S.R.L.

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(54) Title: STABLE LYOPHILIZED PHARMACEUTICAL PREPARATIONS OF MONOCLONAL OR POLYCLONAL
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(57) **Abrégé/Abstract:**

The invention concerns lyophilized pharmaceutical preparations of monoclonal or polyclonal antibodies which contain a sugar or an amino sugar, an amino acid and a surfactant as stabilizers. In addition the invention concerns a process for the production of this stable lyophilisate as well as the use of a sugar or amino sugar, an amino acid and a surfactant as stabilizers for therapeutic or diagnostic agents containing antibodies.



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Abstract

The invention concerns lyophilized pharmaceutical preparations of monoclonal or polyclonal antibodies which contain a sugar or an amino sugar, an amino acid and a surfactant as stabilizers. In addition the invention concerns a process for the production of this stable lyophilisate as well as the use of a sugar or amino sugar, an amino acid and a surfactant as stabilizers for therapeutic or diagnostic agents containing antibodies.

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Boehringer Mannheim GmbH

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**Stable lyophilized pharmaceutical preparations of
monoclonal or polyclonal antibodies**

The invention concerns lyophilized pharmaceutical preparations of monoclonal or polyclonal antibodies which contain a sugar or amino sugar, an amino acid and a surfactant as stabilizer. In addition the invention concerns a process for the production of these stable lyophilisates as well as the use of a sugar or amino sugar, an amino acid and a surfactant as stabilizers of therapeutic or diagnostic agents containing antibodies.

The production of immunoglobulins in particular monoclonal and polyclonal antibodies, for therapeutic and diagnostic purposes is nowadays of major and continuously increasing importance.

The use of antibodies as pharmacological agents has been already known for a long time and comprises numerous applications. Hence antibodies have been for example used successfully for tetanus prophylaxis, to combat pathogenic microorganisms or to neutralize their toxins and also for poisoning by snake venoms.

If the antigen involved in the disease mechanism has been identified, which is the case for numerous infectious and some oncological indications for antibody therapy, one utilizes the specificity of the antibodies for the therapy.

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In clinical and preclinical studies antibodies are presently used to lower the cholesterol level, to influence the angiotensin/renin system and in autoimmune diseases such as for example lupus, autoimmune encephalitis, multiple sclerosis, polyarthrititis and autoimmune myasthenia gravis.

Additionally of major therapeutic importance is their application to counteract intoxications by low molecular substances such as e.g. the Fab fragments of anti-digoxin antibodies when used for intoxications by digoxin or the cardiac glycosides digitoxin and ouabain. Moreover antibodies are used in the diagnostic field to identify, purify and determine the content of proteins.

Genetic engineering which revolutionized the production of monoclonal antibodies in cell cultures in the second half of the 70's and in the 80's has greatly advanced the preparation of antibodies.

In order to fulfil these diverse applications it is necessary to have pharmaceutical preparations of monoclonal and polyclonal antibodies that are stable on storage. There are a number of publications relating to liquid formulations or lyophilisates of special antibodies. Thus for example liquid formulations of antibodies are described in EP 0 280 358, EP 0 170 983, WO 89/11298, EP 0 352 500 and JP 63088197.

According to EP 0 280 358 dextran is added to the antibody solution to stabilize it towards certain hormones by which means it was possible to achieve a stability of over nine months. According to EP 0 170 983 hydrolysed ovalbumin is added to stabilize a

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thermolabile monoclonal antibody when heated and as a result the antibody could still be used after storage at 45°C for 7 days. Polyhydroxy alcohols (e.g. glycerol, inositol, polyvinyl alcohol) or sugars (e.g. sucrose and glucose) or glycitols (e.g. sorbitol, mannitol) are known from JP 63088197 as further stabilizers for liquid formulations. WO 89/11298 demonstrates the use of maltose in a phosphate buffer containing sodium chloride as a further method for the liquid stabilization of monoclonal antibodies. EP 0 352 500 describes polyethylene glycol 4000 and 3-propiolactone for the liquid stabilization of monoclonal antibodies.

However, in general liquid formulations are not an optimal solution due to storage stability since the proteins or aggregates thereof may precipitate in time during storage, at increased temperatures, when transported through different climatic zones or by improper storage (e.g. interruptions in the cool chain) and the solutions may thus have a reduced protein content and become turbid. Hence, a problem-free use of the solutions cannot be guaranteed in these cases.

In contrast in the case of a lyophilisate formulation the removal of water minimizes the formation of degradation products (e.g. by deamidation and hydrolysis) and aggregate formation. The residual content of water (bound water) can contribute to the stability particularly in the presence of sugars (Hsu et al. Dev. Biol. Stand. 1991, 74: 255-267 and Pikal et al., Dev. Biol. Stand. 1991, 74: 21-27).

Lyophilisate formulations with special antibodies as active substances are also known from the literature but

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they do not give consistent advice about the problem of stabilization. Hence in WO 93/00807 the stabilization of biomaterials is described such as human proteins, growth hormones, interleukins, interferons, enzymes and also monoclonal and polyclonal antibodies by a two component system consisting of cryoprotective agents (e.g. polyethylene glycols) and a compound which can form hydrogen bridges with proteins. However, a disadvantage of these preparations is that the addition of high molecular compounds such as polyethylene glycols can lead to an accumulation in the body with potentially toxic side-effects if there is no biodegradation. Furthermore, as is well-known, polymers can act as antigens depending on their molar mass.

Lyophilisates of a monoclonal antibody that is labile when frozen are stabilized for one year according to JP 60146833 by the addition of albumin (human, horse or bovine albumin). Human serum albumin (HSA) is also described in EP 0 303 088 in combination with a carbohydrate (e.g. dextrose, sucrose or maltose) to stabilize a monoclonal antibody for the treatment of *Pseudomonas aeruginosa* infections.

Human serum albumin (in combination with sugars and amino acids) is also the principle by which monoclonal antibodies are stabilized in EP 0 413 188. In JP 01075433 a mixture of human serum albumin, mannitol and polyethylene glycol is used to stabilize a human monoclonal antibody as a lyophilisate. A further example of the use of macromolecules such as e.g. polyethylene glycols and protecting proteins such as human serum albumins to stabilize gamma-globulins during lyophilization is shown in WO 84/00890.

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In WO 93/01835 Hagiwara et al. describe the stabilization of a human monoclonal antibody by lyophilization with mannitol and glycine in a solution containing sodium chloride and phosphate buffer. Stable preparations are obtained with regard to freezing, lyophilization and reconstitution.

Draber et al. (J. Immun. Methods, 1995, 181:37-43) were able to produce a stable formulation of monoclonal IgM antibodies from the mouse at 4°C by the addition of trehalose alone and in combination with polyethylene glycol 8,000. However, the antibodies are only stable for 14 days at 50°C. Using other monosaccharides or disaccharides alone such as e.g. sucrose, maltose, lactose or galactose it is not possible to stabilize these antibodies.

A monoclonal antibody from the mouse is converted into a stable lyophilisate in WO 89/11297 using a carbohydrate (maltose) and a buffer in the acid range (acetate buffer). In this case a disadvantage is the limitation to buffering in an acid range.

Polymeric gelatin as a freezing protectant and stabilizer in a lyophilisate is used in WO 92/15 331. The stabilization is also achieved in combination with a carboxylic acid (e.g. citric acid) or a salt thereof as well as with a primary, secondary or tertiary alcohol or an amino acid in a pH range of 6.8 to 8.1.

In a whole series of the aforementioned publications pharmaceutical additives or auxiliary substances are proposed as stabilizers which are not acceptable from a medical point of view. Hence polymers (such as PEG or

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gelatin) and proteins (such as serum albumins) pose a certain risk due to their origin and their physico-chemical properties and can trigger allergic reactions even to the point of an anaphylactic shock. Proteins of human or animal origin as well as proteins obtained from cell cultures carry the residual risk of viral contaminations. However, other protein-like contaminations which are difficult to detect analytically can cause immunological reactions in humans due to their properties.

The addition of polymeric compounds such as e.g. polyethylene glycols (PEG) or gelatin can lead to an accumulation in the body with potentially toxic side-effects if there is no biodegradation. Polymers may also have antigenic properties depending on their molar mass. Also it is difficult to ensure the purity of polymers due to the catalysts used in their production or the presence of monomers and other polymer fragments. The use of polymers in pharmaceutical forms of administration, especially in drug forms that can be administered subcutaneously, should be avoided if another type of stabilization is possible.

In contrast the use of sugars alone without other additives does not always ensure an adequate protective effect when the antibodies are lyophilized.

Hence the object of the invention was to provide a stable pharmaceutical preparation of monoclonal or polyclonal antibodies that is essentially free of the above-mentioned polymers or proteinaceous pharmaceutical auxiliary substances. This applies particularly to those antibodies which are labile towards freezing and thawing

processes or towards multiple freezing and thawing processes.

According to one aspect of the present invention, there is provided a stable lyophilized pharmaceutical preparation of monoclonal or polyclonal antibodies containing a sugar or an amino sugar, a single basic amino acid, a surfactant, and an inorganic acid, acetic acid or a salt thereof.

According to another aspect of the present invention, there is provided an aqueous pharmaceutical preparation of monoclonal or polyclonal antibodies obtainable by redissolving the lyophilizate as described herein with the proviso that the aqueous pharmaceutical preparation is not obtained by redissolving the lyophilizate in phosphate buffered saline, sterile saline solution or Ringer's solution.

According to still another aspect of the present invention, there is provided a process for the production of an aqueous pharmaceutical preparation of monoclonal or polyclonal antibodies as described herein, wherein the aqueous preparation that is produced contains a monoclonal or polyclonal antibody as the active substance, a sugar or amino sugar, a single basic amino acid, a surfactant and an inorganic acid, acetic acid or a salt thereof as additives, the solution is lyophilized.

According to yet another aspect of the present invention, there is provided the use of a combination of auxiliary substances composed of a sugar or an amino sugar, a single basis amino acid, a surfactant and an inorganic acid, acetic acid or a salt thereof

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for the production of stable lyophilized therapeutic or diagnostic agents containing antibodies.

According to one aspect of the present invention, there is provided a stable lyophilized pharmaceutical preparation of monoclonal antibodies containing trehalose, histidine, polysorbate and histidine hydrochloride.

According to one aspect of the present invention, there is provided an aqueous pharmaceutical preparation of monoclonal or polyclonal antibodies obtainable by redissolving the stable lyophilized pharmaceutical preparation.

According to still another aspect of the present invention, there is provided a process for the production of the stable lyophilized pharmaceutical preparation comprising:

- a) producing an aqueous preparation comprising a monoclonal or polyclonal antibody, a sugar or amino sugar, an amino acid and a surfactant, and
- b) lyophilizing said aqueous preparation.

According to yet another aspect of the present invention, there is provided a stable lyophilized pharmaceutical preparation of monoclonal or polyclonal antibodies, comprising:

- a) sucrose,
- b) a basic amino acid, and
- c) a polyoxyethylene-polyoxypropylene polymer,

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wherein the concentration of the antibodies is up to 20 mg/ml, and wherein said stable lyophilized pharmaceutical preparation does not exhibit any detectable degradation products or protein degradation products or protein aggregates when stored at a temperature of 20 °C for a time of at least 104 weeks.

According to a further aspect of the present invention, there is provided a stable lyophilized pharmaceutical preparation of monoclonal or polyclonal antibodies, comprising:

- a) a sugar or an amino sugar in a concentration of 35-75 mg/ml,
- b) an amino acid in a concentration of 10 mg/ml and
- c) a surfactant in a concentration of 0.01% by weight.

Surprisingly it was found that stable pharmaceutical lyophilisates of monoclonal or polyclonal antibodies are obtained if these contain sugar or amino sugar, an amino acid and a surfactant as additives. The lyophilisates are preferably composed of a) the antibody, b) a sugar or amino sugar, c) an amino acid, d) a buffer for adjusting the pH value and e) a surfactant. Those lyophilisates are particularly preferred which only contain a single or two different amino acids.

These preparations are physiologically well tolerated, have a relatively simple composition and can be dosed exactly. In addition they are stable i.e. they exhibit no detectable degradation products or protein aggregates when subjected to multiple freezing and thawing processes as well as on longer storage. The lyophilisates can even be stored without stability problems at refrigerator temperature (4 - 12°C) or even

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at room temperature (18 - 23°C) over a time period of at least three months, preferably at least six months and in particular of at least one to two years. Furthermore they are also stable when stored at higher temperatures (for example up to 30°C). The storage stability is for example exhibited by the fact that during the said storage period only a very small number of particles can be detected when the lyophilisates are reconstituted in the containers with water for injection purposes or with isotonic solutions. In particular the containers have fewer than 6000 particles with a particle size of more than 10 μm and/or less than 600 particles with a particle size of more than 25 μm . The solutions prepared in this manner are stable over a time period of about up

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to five days, preferably up to three days.

The fact that the preparations protect against freezing due to the selected combination of additives is particularly advantageous. Hence, in particular this enables a lyophilization at temperatures down to -45°C without impairing the stability of the antibodies. In addition the lyophilisates containing the combination of additives according to the invention are also stable for a long period and during storage even at relatively high temperatures. Especially compared to conventional formulations, they exhibit no particle formation after reconstitution with water, i.e. the solutions are essentially free of turbidities.

The preparations according to the invention have the additional advantage of being essentially free of protein-like or polymeric auxiliary substances the use of which may be problematic from a medical point of view. Due to the fact that liquid therapeutic or diagnostic agents containing antibodies with a pH value of about 5 to 8, preferably with a pH value of 6.0 - 7.4 (pH value of blood 7.2 - 7.4) can now be prepared by dissolving lyophilisates, they have the additional advantage of being well-tolerated and can be administered substantially free of pain. This is above all important for subcutaneous administration since in this case intolerances develop more easily than when administered intravenously.

The formulations according to the invention can in general be produced in clinically relevant concentration ranges of the antibody for example of up to 20 mg/ml preferably up to 10 mg/ml. Preferred concentration

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ranges are concentrations above 0.01 mg/ml in particular above 0.05 and 0.1 mg/ml. In particular concentration ranges of 0.05 - 10 mg/ml or 0.1 - 5 mg/ml for example about 5, 8 or 10 mg/ml are used. The injection volumes of the solutions used are less than 2 ml preferably about 1 ml in the case of subcutaneous or intravenous injections. Small injection volumes are particularly advantageous for subcutaneous administration since they only cause slight mechanical irritation in the subcutaneous tissue. Basically the solutions are also directly suitable as additives to infusion solutions or as infusion solutions. If they are used as additives to infusion solutions the concentration of the antibodies is at higher levels, for example up to 10 mg/ml. These concentrated solutions of the antibodies are then added to conventional infusion solutions so that the concentration of the antibody in the infusion solution to be administered is in the therapeutically relevant range. This range is normally 0.001 - 0.5 mg/ml.

The pharmaceutical single forms of administration can either be present as ready-to-use infusion solutions or injection solutions or also as lyophilisates. If the pharmaceutical preparations are used in the form of lyophilisates, the single dose containers, for example glass ampoules with a volume of 10 ml, contain the antibody in amounts of 0.1 - 500 mg, preferably 10 - 100 mg depending on the respective therapeutically relevant dose of the antibody. The lyophilisate optionally contains additional conventional pharmaceutical auxiliary substances. The lyophilisate is dissolved with an appropriate amount of reconstitution solution and can then either be used directly as an injection solution or as an additive to an infusion solution. If it is used as an additive to infusion

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solutions, the lyophilisate is usually dissolved with about 10 ml of a reconstitution solution and added to a physiological saline solution (0.9 % NaCl) of 250 ml. The resulting infusion solution is then usually administered to the patient within about 30 minutes.

The sugars used according to the invention can be monosaccharides, disaccharides or trisaccharides. Glucose, mannose, galactose, fructose and sorbose come into consideration as monosaccharides. Sucrose, lactose, maltose or trehalose come into consideration as disaccharides. Raffinose is preferably used as the trisaccharide. According to the invention sucrose, lactose, maltose, raffinose or trehalose are especially preferably used. Instead of maltose it is also possible to use the stereoisomeric disaccharides cellobiose, gentiobiose or isomaltose.

Those monosaccharides are generally referred to and used as amino sugars which have an amino ($-\text{NH}_2$, $-\text{NHR}$, $-\text{NR}_2$) or an acylated amino group ($-\text{NH}-\text{CO}-\text{R}$) instead of a hydroxy group. For this glucosamine, N-methyl-glucosamine, galactosamine and neuraminic acid are particularly preferred according to the invention. The sugar content or amino sugar content is for example up to 2000 mg, preferably up to 1000 mg especially up to 800 or up to 500 mg per single form of administration. Amounts of more than 10, 50 or 100 mg come for example into consideration as the lower limit for the sugar content. Preferred ranges are 200 - 1000 mg, especially 400 - 800 mg. The stated quantities per single form of administration refer to single forms of administration which are marketed as lyophilisates. Such lyophilisates are preferably filled into injection bottles with a volume of 10 ml. After dissolution of the lyophilisates

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with a reconstitution solution of 10 ml, liquid forms of administration are obtained which can be administered directly. The sugar concentration in these injection solutions is up to 200 mg/ml, preferably up to 100 mg/ml based on the amounts stated above of the sugars used.

The amino acids used according to the invention can be basic amino acids such as arginine, lysine, histidine, ornithine etc., the amino acids preferably being used in the form of inorganic salts thereof (preferably in the form of phosphoric acid salts i.e. as amino acid phosphates). If free amino acids are used, the desired pH value is adjusted by adding a suitable physiologically tolerated buffer substance such as e.g. an inorganic acid in particular phosphoric acid, sulphuric acid, acetic acid, formic acid or salts thereof. In this case the use of phosphates has the particular advantage that particularly stable lyophilisates are obtained. It has proven to be advantageous when the preparations are essentially free of organic acids such as e.g. malic acid, tartaric acid, citric acid, succinic acid, fumaric acid, etc. or the corresponding anions (malates, oxalates, citrates, succinates, fumarates, etc.) are not present.

Preferred amino acids are arginine, lysine or ornithine. In addition it is also possible to use acidic amino acids such as glutamic acid and aspartic acid or neutral amino acids such as e.g. isoleucine, leucine and alanine or aromatic amino acids such as e.g. phenylalanine, tyrosine or tryptophan. The amino acid content in the aqueous preparations according to the invention is up to 100 mg/ml, preferably up to 50 mg/ml or up to 30 mg/ml. The lower limit may for example be concentrations above 1, 5 or 10 mg/ml. Preferred concentrations are for example in the range of 3 - 30 mg/ml or 10 - 125 mg/ml.

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If the corresponding forms of administration are marketed as lyophilisates, these lyophilisates are preferably made available in injection bottles (volumes of for example 10 ml). Such single forms of administration contain the amino acids in amounts of up to 1000 mg, preferably up to 500 mg or up to 300 mg.

Surfactants which come into consideration are all surfactants that are usually used in pharmaceutical preparations preferably polysorbates and polyoxy-ethylene-polyoxypropylene polymers such as e.g. Tween®. Low amounts of surfactant of 0.05 to 0.5 mg/ml preferably 0.1 mg/ml are sufficient to stabilize the antibodies. In the above-mentioned single forms of administration the amount of surfactants is 0.5 - 5 mg in the case of a lyophilisate that is filled into an injection bottle of 10 ml.

The stabilization of antibodies achieved by the said additives relates in principle to all known monoclonal and polyclonal antibodies and their Fab fragments. Humanized antibodies and modified antibodies (cf. e.g. US 5,624,821; EP 0 592 106; PCT/EP96/00098) are preferably used. The molecular weight of the antibodies is 50 kDa-200 kDa per monomer unit, in particular the molecular weight is about 80 - 150 kDa. In particular antibodies to the hepatitis B virus (cf. WO 94/11495), to AIDS viruses, cytomegalo viruses, meningoencephalitis viruses (FSME), rubella viruses, measles viruses, rabies pathogens, *Pseudomonas aeruginosa* bacteria, varicella-zoster viruses, tetanus pathogens, van Willebrandt factor (cf. WO 96/17078), NGFR (nerve growth factor receptor), PDGFR (platelet derived growth factor receptor: Shulman, Sauer, Jackman, Chang, Landolfi, J. Biol. Chem. 1997, 272(28): 17400-4), selectin, in particular E-selectin,

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L-selectin (cf. Takashi et al., Proc. Natl. Acad. Sci. USA 1990, 87: 2244-2248; WO 94/12215) or P-selectin; integrins or diphtheria pathogens can be stabilized according to the invention. The antibody concentration can preferably be up to 8 mg/ml. It is preferably for example 0.05 - 2 mg/ml. The amount of antibody in the single form of administration, for example in a lyophilisate in an injection bottle of 10 ml, is up to 100 mg preferably up to 80 mg, 50 mg, 20 mg or 10 mg. The concentration of the antibodies after reconstitution of the lyophilisates with a volume of 10 ml is in the range of 1 - 10 mg/ml, preferably at 5 - 8 mg/ml.

In addition to the said additives, sugar, amino acid and surfactant, the lyophilisates according to the invention can contain physiologically tolerated auxiliary substances from the group comprising acids, bases, buffers or isotonizing agents to adjust the pH value to 5 to 8, preferably 6.0 to 7.4. The buffer capacity of the preparations is adjusted such that when the lyophilisates are dissolved with standard reconstitution solutions such as for example water for injection purposes the buffer concentration is in the range between 10 - 20 mmol/l preferably at about 15 mmol/l.

The order of addition of the various auxiliary substances or of the antibody is largely independent of the production process and is up to the judgement of a person skilled in the art. The desired pH value of the solution is adjusted by adding bases such as for example alkali hydroxides, alkaline earth hydroxides or ammonium hydroxide. Sodium hydroxide is preferably used for this. The desired pH value can in principle be adjusted by adding basic solutions. In this sense salts of strong bases with weak acids are generally suitable such as

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sodium acetate, sodium citrate, di-sodium or sodium dihydrogen phosphate or sodium carbonate. If the pharmaceutical solution of auxiliary substances has a basic pH value it is adjusted by titration with an acid until the desired pH range has been reached.

Physiologically tolerated inorganic or organic acids come into consideration as acids such as for example hydrochloric acid, phosphoric acid, acetic acid, citric acid or conventional solutions of substances which have an acidic pH value. In this sense preferred substances are salts of strong acids with weak bases such as e.g. sodium dihydrogen phosphate or disodium hydrogen phosphate. The pH value of the solution is preferably adjusted with phosphoric acid or an aqueous sodium hydroxide solution.

In order to produce well-tolerated parenteral drug forms it is expedient to add isotonizing auxiliary substances if isotonicity cannot be already achieved by the osmotic properties of the antibody and the additives used for stabilization. Non-ionized well-tolerated auxiliary substances are used above all for this. Salts such as NaCl should, however, only be added in small amounts, in particular a value of 30 mmol/l in the final injection or infusion solution for administration should not be exceeded.

In addition the pharmaceutical preparations can contain further common auxiliary substances or additives. Antioxidants such as for example glutathione or ascorbic acid or similar substances can be added.

For the production of the lyophilisates the aqueous pharmaceutical solutions which contain the antibody are

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firstly produced. A buffered antibody solution containing sodium chloride is preferably prepared. This antibody solution is admixed with an aqueous solution containing the additives sugar, amino acid and surfactant during which the pH value is adjusted with an acid or base to 5 to 8. Phosphoric acid or phosphate salts and sodium chloride are added in such amounts that the previously defined concentrations are obtained. Subsequently it is sterilized by filtration and the solution prepared in this manner is lyophilized.

The invention also enables unstable aqueous solutions containing antibodies that are sensitive to freezing to be also converted by means of freeze-drying into stable preparations that are also stable at high temperatures without impairing the quality.

A further advantage of the lyophilisates according to the invention is that, in addition to avoiding damage to the antibodies during freezing, they also exhibit no reduction in the antibody content and no aggregate formation or flocculation even after a long-term storage at 50°C. They are thus stable with regard to antibody content and purity. The formation of particles is prevented which is exhibited by the low values for turbidity after reconstitution of the lyophilisates with water for injection purposes.

The invention is elucidated in more detail in the following on the basis of examples of application.

Examples 1 to 10 show in which manner the lyophilisates according to the invention can be formulated, produced and examined with regard to antibody stability.

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Comparative experiments without auxiliary substances or with sucrose alone or with mannitol as a substitute for the sugar component or with the amino acid component alone or only the sugar or amino acid component without the surfactant show that the choice of the combination of additives according to the invention is essential for achieving a stable formulation. Sucrose alone, amino acid alone or both components without surfactant lead to unstable formulations.

The formulations according to the invention are insensitive to freezing and it is possible to completely omit polymers or proteins that are regarded as being toxic such as polyethylene glycols, gelatin, serum albumins. In the case of the surfactants only relatively small amounts of physiologically well-tolerated surfactants are present.

The antibody to HBV used in the following application examples is a recombinant human monoclonal antibody (MAB) from a murine cell. It has a molecular weight of about 147 kDa and is directed towards the hepatitis B surface antigen (HBsAg) of the hepatitis B virus. The monoclonal antibody recognizes the a-determinant of the HBsAg which is constant in almost all known variants of the virus. This antibody can for example be used for the following medical indications: treatment of chronic hepatitis for which there has previously not yet been a satisfactory treatment method; treatment of passive immunoprophylaxis in HBsAg-positive liver transplant patients. In central and northern Europe and the USA up to 2 % of the population are carriers of the hepatitis B virus, in southern Europe up to 3 %, in Africa and the Far East it is 10 - 15 %. A consequence of this chronic infection is that the risk of developing hepatocellular

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carcinoma is increased by 100-fold, 40 % of the virus carriers die as a result of this infection.

Antibodies to L-selectin, the NGF receptor or the PDGF receptor can be preferably used as antibodies within the sense of the invention.

Example 1 shows the properties of an aqueous solution of a monoclonal antibody to hepatitis B virus (MAB HBV; INN name: Tuvirumab) containing phosphate buffer and sodium chloride at pH = 5, pH = 6.5 and pH = 8 after freezing and thawing. It shows that freezing and thawing damages the monoclonal antibody.

Example 2 demonstrates the possibility of stabilizing a preparation according to the invention with sucrose or maltose or an amino sugar (N-methylglucosamine or galactosamine) and arginine phosphate and Tween 20 with a concentration of the antibody of 2 mg/ml i.e. 2 mg in the lyophilisate.

The same preparation as in example 2 is shown in example 2a except that the antibody concentration is 8 mg/ml. It can be seen from examples 2 and 2a that the combination of the said auxiliary substances not only avoids damage to the antibody during freezing but also has a positive influence on the stability during long-term storage.

Example 3 elucidates the necessity of amino acids and surfactant in the preparation according to the invention. The use of sucrose as a builder alone leads to an unstable lyophilisate.

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Example 4 describes variations of the amino acid component. It turns out that variation of the basic amino acids in the form of arginine or ornithine as well as the substitution of the basic amino acid by a neutral amino acid such as e.g. by leucine or by an acidic amino acid such as e.g. aspartic acid leads to a storage-stable preparation.

In example 5 the lyophilisation of a formulation containing sucrose, arginine and Tween 20 as well as phosphate buffer and sodium chloride is compared at various pH values (pH 5, pH 6.5 and pH 8). The data obtained show that it is possible to lyophilize within this pH range without impairing the stability.

If the said surfactant Tween 20 is replaced by a representative of the surface-active class of compounds polyoxyethylene-polyoxypropylene polymers (commercial name Pluronic[®]) as in example 6 this also results in an adequate stability of the preparation according to the invention.

Example 7 demonstrates the instability of a formulation containing mannitol as the builder as a substitute for sucrose, maltose or the amino sugar (see example 2).

If the sugar and the surfactant are omitted in the formulation the preparation becomes unstable as shown in example 8.

Although a combination of sugar (e.g. sucrose) and amino acids without surfactant in example 9 yields good results with regard to the parameters content and aggregates, the turbidity is, however, substantially

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increased compared to the formulations according to the invention containing sugar, amino acid and surfactant.

Example 10 shows that other monoclonal antibodies can also be stabilized with a combination of the sugar, amino acid and surfactant. The antibody anti-L-selectin is for example stable at a concentration of 7 mg in the lyophilisate. The lyophilization is carried out starting with a volume of 1 ml of an aqueous solution.

Investigative methods to determine stability

The lyophilized preparations were stored under defined storage conditions in the absence of light and subsequently analysed. The following test methods were used for the analyses.

OD280: Optical density at 280 nm. Photometric determination of the protein content, the UV absorbance is due to side chain chromophores such as tryptophan tyrosine and phenylalanine residues. Specification: 95-105 %.

SE-HPLC: Size-exclusion high performance chromatography to determine aggregates. Specification: max. 2 %.

Measurement of turbidity: After reconstituting the lyophilisate the undiluted antibody solution was measured in a suitable turbidity photometer. Specification: max. 6 turbidity units.

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Example 1:

An aqueous stock solution of the MAB to HBV described above containing phosphate buffer and sodium chloride is prepared and examined. The concentration of the MAB is about 15 mg/ml.

Table 1a shows on the one hand the lability to freezing of the monoclonal antibody solution at various pH values at -20°C which already results in a decrease of the protein content after 4 weeks to 92.1 and 94.2 and 94.0 %. A decrease of the protein content is also observed on storage at 25°C. Under the storage conditions 4-8°C in a refrigerator the antibody is adequately stable over 9 months.

Tables 1b-1d show the stability data of the monoclonal antibody solution prepared at pH values 5, 6.5 and 8 at -20°C, 4-8°C and 25°C. This also shows that only a storage at 4-8°C is acceptable.

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Table 1a: Change of the antibody content in the solution of active substance (10 mM phosphate buffer, 30 mM sodium chloride, water for injection purposes)

	pH 5			pH 6.5			pH 8		
Time	-20°C	4-8°C	25°C	-20°C	4-8°C	25°C	-20°C	4-8°C	25°C
start		> 99			> 99			> 99	
4 weeks	92.1	> 99	> 99	94.2	> 99	> 99	94.0	> 99	> 99
13 weeks	78.9	> 99	97.2	81.2	> 99	98.1	77.8	> 99	96.1
6 months	61.2	> 99	94.1	69.9	> 99	94.4	65.8	> 99	91.9
9 months	47.8	> 99	88.7	55.6	> 99	90.2	51.0	> 99	84.3

All data in %. The protein was determined by measuring the absorbance at 280 nm (OD 280).

Table 1b: Aggregate formation and turbidity values for the active substance solution of antibody, pH = 5

Times	-20°C aggregates turbidity		4-8°C aggregates turbidity		25°C aggregates turbidity	
start	n.d.	1.5	n.d.	1.5	n.d.	1.5
4 weeks	aggregates	floccul.	n.d.	1.5	0.7 %	1.5
13 weeks	aggregates	floccul.	0.2 %	1.8	1.9 %	1.8
6 months	aggregates	floccul.	0.3 %	1.9	aggregates	9.9
9 months	aggregates	floccul.	0.6 %	2.1	aggregates	10.9

n.d.= not detectable

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Table 1c): Aggregate formation and turbidity values for the active substance solution of antibody, pH = 6.5

Times	-20°C aggregates turbidity	4-8°C aggregates turbidity	25°C aggregates turbidity
start	n.d. 1.2	n.d. 1.2	n.d. 1.2
4 weeks	aggregates floccul.	n.d. 1.3	0.5 % 1.4
13 weeks	aggregates floccul.	0.2 % 1.4	1.8 % 1.7
6 months	aggregates floccul.	0.3 % 1.9	4.9 % floccul.
9 months	aggregates floccul.	0.6 % 2.1	9.3 % floccul.

Table 1d: Aggregate formation and turbidity values for the active substance solution of antibody, pH = 8

Times	-20°C aggregates turbidity	4-8°C aggregates turbidity	25°C aggregates turbidity
start	n.d. 1.4	n.d. 1.4	n.d. 1.4
4 weeks	2.0 % floccul.	0.3 % 1.5	0.74 % 1.7
13 weeks	2.8 % floccul.	0.5 % 1.8	1.95 % 2.1
6 months	3.7 % floccul.	0.6 % 1.9	3.0 % floccul.
9 months	5.4 % floccul.	0.8 % 2.1	4.3 % floccul.

Aggregates in % using SE-HPLC, turbidity in turbidity units (turbidity) using a turbidity photometer.

Example 2:

A solution of the monoclonal antibody to HBV according to example 1 was added to aqueous solutions of the

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following sugars or amino sugars: sucrose (formulation 1), maltose (formulation 2) and N-methylglucosamine (formulation 3) containing arginine phosphate buffer and Tween 20 as the surfactant. The formulation is listed in example 2a. The final concentration of the MAB is 2 mg/ml. After adjusting the pH value with phosphoric acid to 6.5, the solutions were sterilized by filtration (0.22 μ m membrane filter) and filled into sterilized and depyrogenized injection bottles made of glass (hydrolytic class I) (filling volumes 1 ml) and lyophilized. After lyophilization the injection bottles were aerated with nitrogen, sealed automatically with stoppers in the freeze drying chamber and subsequently flanged.

The flanged injection bottles were stored in the absence of light for 4 to 13 weeks at various temperatures. After these periods the stability of the lyophilisates was examined with the described methods of examination.

Table 2a: Storage at 25°C

	Storage 4 weeks at 25°C			Storage 13 weeks at 25°C		
	I	II	III	I	II	III
form. 1 sucrose	100	n.d.	1.7	100	n.d.	1.6
form. 2 maltose	100	n.d.	1.6	100	n.d.	1.8
form. 3 N-methyl-glucosamine	100	n.d.	1.8	100	n.d.	1.5

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Table 2b: Storage at 50°C

	Storage 4 weeks at 50°C			Storage 13 weeks at 50°C		
	I	II	III	I	II	III
form. 1 sucrose	>99	n.d.	2.0	>99	n.d.	2.0
form. 2 maltose	>99	n.d.	1.9	>99	n.d.	2.1
form. 3 N-methyl-glucosamine	>99	n.d.	1.7	>99	n.d.	2.0

Legend:

I protein content in % with OD 280

II aggregates in % with SE-HPLC

III turbidity of the reconstituted solution in
turbidity units (dimensionless number)n.d. not detectable (used in the same way in all further
tables)**Example 2a**

In example 2a the formulation 1 from example 2 is prepared with 8 mg/ml antibody (= formulation 1a). It turns out that higher concentrations of up to 8 mg/1 ml antibody are adequately stable in this formulation.

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Compositions of formulations 1 and 1a:

	Formulation 1	Formulation 1a
MAB HBV	2.0 mg	8.0 mg
phosphate buffer	15 mM	15 mM
sodium chloride	30 mM	30 mM
sucrose	68.0 mg	58.0 mg
arginine	10.0 mg	10.0 mg
phosphoric acid	ad pH 6.5	ad pH 6.5
Tween 20	0.1 mg	0.1 mg
water for injection purposes	ad 1.0 ml	ad 1.0 ml

Table 3a: Stability data for formulation 1 and formulation 1a at 25°C

	Storage 4 weeks at 25°C			Storage 13 weeks at 25°C		
	I	II	III	I	II	III
form.1: 2 mg/1ml	100	n.d.	1.7	100	n.d.	1.6
form.1a:8 mg/1ml	>99	n.d.	4.8	>99	n.d.	4.7

Table 3b: Stability data for formulations 1 and 1a at 50°C

	Storage 4 weeks at 50°C			Storage 13 weeks at 50°C		
	I	II	III	I	II	III
form.1: 2 mg/1ml	>99	n.d.	2.0	>99	n.d.	2.0
form.1a:8 mg/1ml	>99	n.d.	4.7	>99	n.d.	5.5

I protein content in % with OD 280

II aggregates in % with SE-HPLC

III turbidity of the reconstituted solution in
turbidity units (dimensionless number)

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Example 3

Comparison of formulations 1 and 4. Formulation 4 only contains sucrose as the builder and no arginine phosphate and no Tween 20. Formulation 4 is unstable.

	Formulation 1	Formulation 4
MAB HBV	2.0 mg	2.0 mg
phosphate buffer	15 mM	15 mM
sodium chloride	30 mM	30 mM
sucrose	68.0 mg	68.0 mg
arginine	10.0 mg	--
phosphoric acid or NaOH	ad pH 6.5	ad pH 6.5
Tween 20	0.1 mg	--
water for injection purposes	ad 1.0 ml	ad 1.0 ml

Table 4a: Storage at 25°C

	Storage 4 weeks at 25°C			Storage 13 weeks at 25°C		
	I	II	III	I	II	III
form.1: sucrose with arg. phos. and Tween 20	100	n.d.	1.7	>99	n.d.	1.6
form.4: sucrose without arg.phos. and Tween 20	98.3	1.6	6.1	96.0	4.3	9.5

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Table 4b: Storage at 50°C:

	Storage 4 weeks at 50°C			Storage 13 weeks at 50°C		
	I	II	III	I	II	III
form.1:sucrose with arg.phos. and Tween 20	100	n.d.	2.0	>99	n.d.	2.0
form.4:sucrose without arg.phos. and Tween 20	96.0	4.2	8.5	89.8	10.1	10.9

Legend:

- I protein content in % with OD 280
- II aggregates in % with SE-HPLC turbidity of the reconstituted solution in turbidity units (dimensionless number)
- III turbidity of the reconstituted solution in turbidity units (dimensionless number)

Example 4

Variation of the amino acid component of the formulation. Formulations with basic, acidic and neutral amino acids are stable.

Composition of the formulations:

MAB HBV	2.0 mg
phosphate buffer	15 mM
sodium chloride	30 mM
sucrose	35 - 70 mg
amino acid	variable
phosphoric acid or NaOH	ad pH 6.5
Tween 20	0.1 mg
water for injection purposes	ad 1.0 ml

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Table 5

	Amino acid
formulation 1	arginine (basic)
formulation 5	ornithine (basic)
formulation 6	leucine (neutral)
formulation 7	aspartic acid (acidic)

The pH value is adjusted by phosphoric acid or hydroxide solution.

Tables 6a - d

Examination results of formulations 1, 5, 6, 7 after storage for 4 and 13 weeks.

a) Table 6a, Formulation 1 (arginine):

	Storage period 25°C	4 weeks 50°C	Storage period 25°C	13 weeks 50°C
protein content % (OD 280)	100	>99	100	>99
aggregates % (SE-HPLC)	n.d.	n.d.	n.d.	n.d.
turbidity	1.7	2.0	1.6	2.0

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b) Table 6b, formulation 5 (ornithine):

	Storage period 25°C	4 weeks 50°C	Storage period 25°C	13 weeks 50°C
protein content % (OD 280)	>99	>98	>98	>98
aggregates % (SE-HPLC)	n.d.	n.d.	n.d.	n.d.
turbidity	1.9	1.9	2.0	2.1

c) Table 6c, formulation 6 (leucine):

	Storage period 25°C	4 weeks 50°C	Storage period 25°C	13 weeks 50°C
protein content % (OD 280)	>98	>98	>98	>98
aggregates % (SE-HPLC)	n.d.	n.d.	0.1	0.1
turbidity	2.2	2.4	2.8	2.7

d) Table 6d, formulation 7 (aspartic acid):

	Storage period 25°C	4 weeks 50°C	Storage period 25°C	13 weeks 50°C
protein content % (OD 280)	>98	>98	>98	>98
aggregates % (SE-HPLC)	n.d.	n.d.	0.1	0.1
turbidity	2.7	2.7	3.4	4.0

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Example 5

Example 5 contains formulation 1 at various pH values, the lyophilisates are prepared as described in example 2, the pH of the solution of auxiliary substances and of the product solution was adjusted before lyophilisation with 85 % phosphoric acid.

Formulation:

MAB HBV	2.0 mg
phosphate buffer	15 mM
sodium chloride	30 mM
sucrose	68 mg
arginine	10 mg
phosphoric acid	ad pH 5; 6.5; 8
Tween 20	0.1 mg
water for injection purposes	ad 1.0 ml

The lyophilisates were prepared with the pH values shown in table 7.

After flanging the injection bottles these were stored in the absence of light under defined temperature conditions. After storage periods of 4 weeks and 13 weeks the samples were analysed (protein content in %: OD 280, aggregates in %: SE-HPLC, turbidity). The formulation was stable at all pH values.

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Table 7:

	pH
formulation 8	5
formulation 9 (ident. with 1)	6.5
formulation 10	8

Table 8a: Storage at 25°C

	Storage 4 weeks at 25°C			Storage 13 weeks at 25°C		
	I	II	III	I	II	III
formulation 8	100	n.d.	1.9	>99	n.d.	2.3
formulation 9(=1)	100	n.d.	1.7	100	n.d.	1.6
formulation 10	>99	n.d.	2.3	>99	n.d.	2.6

Table 8b: Storage at 50°C

	Storage 13 weeks at 50°C			Storage 13 weeks at 50°C		
	I	II	III	I	II	III
formulation 8	>99	n.d.	2.2	>99	n.d.	2.3
formulation 9(=1)	>99	n.d.	2.0	>99	n.d.	2.0
formulation 10	>98	n.d.	2.5	>98	n.d.	2.6

Legend:

- I protein content in % with OD 280
 II aggregates in % with SE-HPLC
 III turbidity of the reconstituted solution in
 turbidity units (dimensionless number)

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Example 6

The formulation described in the following containing the surfactant Pluronic F 68 instead of Tween 20 was prepared as described above.

The storage and examination of stability was carried out in an analogous manner to that of the other examples.

Formulation 11:

MAB HBV	2.0 mg
phosphate buffer	15 mM
sodium chloride	30 mM
sucrose	48.0 mg
arginine	10.0 mg
phosphoric acid	ad pH 6.5
Pluronic F 68	0.1 mg
water for injection purposes	ad 1.0 ml

Formulation 1 is chosen as a comparison and is identical to formulation 11 except for Tween 20 instead of Pluronic F 68. Both formulations were stable.

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Table 9: Stability data of the formulation containing the surfactants Pluronic F 68 and Tween 20.

	formulation 11				formulation 1			
	storage period		storage period		storage period		storage period	
	4 weeks		13 weeks		4 weeks		13 weeks	
	25°C	50°C	25°C	50°C	25°C	50°C	25°C	50°C
protein content % (OD 280)	>98	>98	>98	>98	100	>99	100	>99
aggregates % (SE-HPLC)	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
turbidity	1.9	1.9	2.5	2.2	1.7	2.0	1.6	2.0

Example 7:

The formulation 12 described in this example essentially corresponds to formulation 1 except that mannitol was used instead of sucrose as a builder. It can be seen that the mannitol formulation is unstable.

Formulation 12:

MAB HBV	2.0 mg
phosphate buffer	15 mM
sodium chloride	30 mM
mannitol	25.0 mg
arginine	10.0 mg
phosphoric acid	ad pH 6.5
Tween 20	0.1 mg
water for injection purposes	ad 1.0 ml

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Table 10: Stability data of the formulations containing the builder mannitol (formulation 12) and sucrose (formulation 1)

	formulation 12				formulation 1			
	storage period		storage period		storage period		storage period	
	4 weeks		13 weeks		4 weeks		13 weeks	
	25°C	50°C	25°C	50°C	25°C	50°C	25°C	50°C
protein content % (OD 280)	96.2	91.8	94.0	84.5	100	>99	100	>99
aggregates % (SE-HPLC)	3.6	8.4	5.8	15.9	n.d.	n.d.	n.d.	n.d.
turbidity	3.2	6.9	4.9	13.2	1.7	2.0	1.6	2.0

Example 8

Further evidence for the necessity of the combination of sugar, amino acid and surfactant is given by comparing formulation 1 which contains all listed components with formulation 13 composed of antibody, phosphate buffer, sodium chloride and arginine phosphate. The aggregate formation is increased and the turbidity values become worse without sugar and surfactant.

Formulation 13

MAB HBV	2.0 mg
phosphate buffer	15 mM
sodium chloride	30 mM
arginine	35.0 mg
phosphoric acid	ad pH 6.5
water for injection purposes	ad 1.0 ml

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Table 11: Stability data for formulation 13 (without sucrose and Tween 20 only with arginine phosphate as builder) and formulation 1

	formulation 13				formulation 1			
	storage period		storage period		storage period		storage period	
	4 weeks		13 weeks		4 weeks		13 weeks	
	25°C	50°C	25°C	50°C	25°C	50°C	25°C	50°C
protein content % (OD 280)	97.6	94.9	95.8	89.0	100	>99	100	>99
aggregates % (SE-HPLC)	2.6	4.5	4.0	10.7	n.d.	n.d.	n.d.	n.d.
turbidity	2.9	4.5	3.8	12.3	1.7	2.0	1.6	2.0

Example 9

Although a stable formulation is obtained without surfactant (Tween 20) and only with sucrose and arginine, the turbidity values worse (formulation 14).

Formulation 15:

MAB HBV	2.0 mg
phosphate buffer	15 mM
sodium chloride	30 mM
sucrose	68.0 mg
arginine	10.0 mg
phosphoric acid	ad pH 6.5
water for injection purposes	ad 1.0 ml

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Table 12: Stability data of formulation 14 and formulation 1

	formulation 14				formulation 1			
	storage period		storage period		storage period		storage period	
	4 weeks		13 weeks		4 weeks		13 weeks	
	25°C	50°C	25°C	50°C	25°C	50°C	25°C	50°C
protein content % (OD 280)	>99	>98	>98	>98	100	>99	100	>99
aggregates % (SE-HPLC)	0.2	0.3	0.5	1.3	n.d.	n.d.	n.d.	n.d.
turbidity	3.4	4.8	8.8	13.3	1.7	2.0	1.6	2.0

Example 10

The following table shows the components of formulation 15. The antibody used is anti-L-selectin. The data shown in table 13a on the examination of stability show that the formulation used enables an adequate stabilization.

Composition of formulation 15:

	Formulation 15
anti-L-selectin	7.0 mg
phosphate buffer	15 mM
sodium chloride	30 mM
sucrose	68.0 mg
arginine	10.0 mg
phosphoric acid	ad pH 6.5
Tween 20	0.1 mg
water for injection purposes	ad 1.0 ml

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Table 13a: Stability data for formulation 15 at 25°C

	Storage 4 weeks at 25°C			Storage 13 weeks at 25°C		
	I	II	III	I	II	III
form. 15:7 mg/1ml	>99	n.d.	2.5	>99	n.d.	2.9

Table 13b: Stability data for formulation 15 at 50°C

	Storage 4 weeks at 50°C			Storage 13 weeks at 50°C		
	I	II	III	I	II	III
form. 15:7 mg/ml	99	n.d.	4.1	99	n.d.	5.2

I protein content in % with OD 280
 II aggregates in % with SE-HPLC
 III turbidity of the reconstituted solution in
 turbidity units (dimensionless number)

Example 11

Stabilization of the antibody anti-L-NGFR (anti-L-nerve-
 growth-factor-receptor)

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Formulation 16:

A lyophilisate with the following formulation (analogous to formulation 1) is prepared:

	Formulation 16
anti-L-NGFR	0.25 mg
phosphate buffer	15 mM
sucrose	75 mg
arginine	10 mg
phosphoric acid	ad pH 6.5
Tween 20	0.1 mg
water for injection purposes	ad 1.0 ml

The lyophilisate of anti-L-NGFR is prepared analogously to the preparation of the MAB-HBV and anti-L-selectin lyophilisates.

An aqueous solution containing the additives sugar, amino acid and surfactant at pH 5 to 8 is admixed with a solution of the anti-L-NGFR in a phosphate buffer. The phosphate salts are added in such amounts that the previously defined concentrations are obtained. Subsequently it is sterilized by filtration and the solution prepared in this manner is lyophilized. After lyophilisation one obtains an optically perfect lyophilisation cake. The antibody anti-L-NGFR remains stable. After reconstitution of the lyophilisate with water for injection purposes a clear solution is obtained.

Claims:

1. A stable lyophilized pharmaceutical preparation of monoclonal or polyclonal antibodies, obtained by combining and lyophilizing:
 - a) a sucrose solution in a concentration of 35 mg/ml to 75 mg/ml;
 - b) an amino acid solution in a concentration of 10 mg/ml, wherein the amino acid is selected from the group consisting of arginine, ornithine, isoleucine and aspartic acid;
 - c) a surfactant solution in a concentration of 0.1 mg/ml, wherein the surfactant is a polysorbate or a polyoxyethylene-polyoxypropylene polymer; and
 - d) monoclonal or polyclonal antibodies.
2. The preparation according to claim 1, wherein the preparation is essentially free of polyethylene glycols.
3. The preparation according to claim 1 or 2, wherein the preparation is essentially free of protein-like standard pharmaceutical auxiliary substances.
4. The preparation according to any one of claims 1 to 3, further comprising physiologically tolerated auxiliary substances selected from the group consisting of acids, bases, buffers and isotonizing agents.
5. The preparation according to any one of claims 1 to 4, wherein the concentration of the antibodies is between 0.25 mg/ml to 8 mg/ml, prior to lyophilisation.
6. An aqueous pharmaceutical preparation of monoclonal or polyclonal antibodies obtainable by redissolving the stable lyophilized pharmaceutical preparation according to any one of claims 1 to 5.

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7. The aqueous pharmaceutical preparation according to claim 6, wherein the solution has a pH value of 5 to 8.
8. The aqueous pharmaceutical preparation according to claim 6, wherein the solution has a pH value of 6 to 7.4.
9. A process for the production of the stable lyophilized pharmaceutical preparation according to any one of claims 1 to 5, comprising:
 - a) producing an aqueous preparation comprising sucrose in a concentration of 35 mg/ml to 75 mg/ml; an amino acid in a concentration of 10 mg/ml, wherein the amino acid is selected from the group consisting of arginine, ornithine, isoleucine and aspartic acid; and a surfactant in a concentration of 0.1 mg/ml, wherein the surfactant is a polysorbate or a polyoxyethylene-polyoxypropylene polymer; and
 - b) lyophilizing said aqueous preparation.
10. The process according to claim 9, further comprising adding pharmaceutical auxiliary substances to the aqueous preparation.
11. The process according to claim 9 or 10, wherein the monoclonal or polyclonal antibody is present in a concentration of 0.25 mg/ml to 8 mg/ml prior to lyophilization.
12. The stable lyophilized pharmaceutical preparation according to any one of claims 1 to 5, wherein said stable lyophilized pharmaceutical preparation does not exhibit any detectable degradation products or protein aggregates when stored at a temperature of 18 to 23°C for a time period of at least 1 to 2 years.
13. The stable lyophilized pharmaceutical preparation according to claim 12, wherein said preparation does not exhibit any detectable degradation products or protein

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aggregates when subjected to multiple freezing and thawing processes.

14. The stable lyophilized pharmaceutical preparation according to claim 12 or 13, wherein the concentration of the antibodies is at least 0.25 mg/ml, prior to lyophilisation.
15. The stable lyophilized pharmaceutical preparation according to claim 12 or 13, wherein the concentration of the antibodies is at least 2.0 mg/ml, prior to lyophilisation.
16. The stable lyophilized pharmaceutical preparation according to claim 12 or 13, wherein the concentration of the antibodies is at least 7.0 mg/ml, prior to lyophilisation.
17. The stable lyophilized pharmaceutical preparation according to claim 12 or 13, wherein the concentration of the antibodies is at least 8.0 mg/ml, prior to lyophilisation.