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STABILIZED TERIPARATIDE SOLUTIONS

TECHNICAL FIELD

This invention relates to pharmaceutical compositions containing a
5 parathyroid hormone. More particularly, the invention relates to teriparatide,
PTH(1-34), stabilized solution formulations

BACKGROUND OF THE INVENTION

Parathyroid hormone (PTH) is a secreted, 84 amino acid product of the
10 mammalian parathyroid gland that controls serum calcium levels through its action
on various tissues, including bone. Studies in humans with certain forms of PTH
have demonstrated an anabolic effect on bone, and have prompted significant
interest in its use for the treatment of osteoporosis and related bone disorders.

Using the N-terminal 34 amino acids of the bovine and human hormone for
15 example, which by all published accounts are deemed biologically equivalent to the
full length hormone, it has been demonstrated in humans that parathyroid hormone
enhances bone growth particularly when administered in pulsatile fashion by the
subcutaneous route. A slightly different form of PTH, human PTH(1-38) has shown
similar results.

20 PTH preparations have been reconstituted from fresh or lyophilized
hormone, and incorporate various forms of carrier, excipient and vehicle. Most are
prepared in water-based vehicles such as saline, or water acidified typically with
acetic acid to solubilize the hormone. The majority of reported formulations also
incorporate albumin as a stabilizer (see for example Reeve et al., Br. Med. J., 1980,
25 280:6228; Reeve et al., Lancet, 1976, 1:1035; Reeve et al., Calcif. Tissue Res., 1976,
21:469; Hodsman et al., Bone Miner; 1990, 9(2):137; Tsai et al., J. Clin. Endocrinol
Metab., 1989, 69(5):1024; Isaac et al., Horm. Metab. Res., 1980, 12(9):487; Law et
al., J. Clin Invest. 1983, 72(3):1106; and Hulter, J. Clin Hypertens, 1986, 2(4):360).
Other reported formulations have incorporated an excipient such as mannitol, which
30 is present either with the lyophilized hormone or in the reconstitution vehicle.
Formulations representative of those employed for human studies include a human



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PTH(1-34) (SEQ ID NO: 2) preparation consisting, upon reconstitution, of mannitol, heat inactivated human serum albumin, and caproic acid (a protease inhibitor) as absorption enhancer (see Reeve et al., 1976, *Calcif. Tissue Res.*, 21, Suppl., 469-477); a human PTH(1-38) preparation reconstituted into a saline vehicle (see Hodsman et al., 1991, 14(1), 67-83); and a bovine PTH(1-34) preparation in aqueous vehicle pH adjusted with acetic acid and containing albumin. There is also an International Reference preparation which for human PTH (1-84) (SEQ ID NO: 1) consists of 100 ng of hormone ampouled with 250 µg human serum albumin and 1.25 mg lactose (1981), and for bovine PTH (1-84) consists of 10 µg lyophilized hormone in 0.01M acetic acid and 0.1% w/v mannitol (see Martindale, *The Extra Pharmacopoeia*, The Pharmaceutical Press, London, 29th Edition, 1989 at p. 1338).

A recent attempt at improving the stability for the lyophilized preparation of h-PTH(1-34) (SEQ ID NO: 2) is reported in EP 619 119 with a combination of sugar and sodium chloride. Also U.S. Pat. No. 5,496,801 describes a freeze-dried composition for the natural hormone, PTH(1-84), containing mannitol as an excipient and a citrate source as a non-volatile buffering agent.

Commercial exploitation of parathyroid hormone requires the development of a formulation that is acceptable in terms of storage stability and ease of preparation. Because it is a protein and thus far more labile than the traditionally small molecular weight drugs, however, the formulation of parathyroid hormone presents challenges not commonly encountered by the pharmaceutical industry. Furthermore, like other proteins that have been formulated successfully, PTH is particularly sensitive to oxidation, deamidation and hydrolysis, and requires that its N-terminal and C-terminal sequences remain intact in order to preserve bioactivity.

It is an object of the present invention to provide a pharmaceutically useful PTH preparation, particularly one comprising, as active ingredient, teriparatide, PTH(1-34) (SEQ ID NO: 2).

CLAIMS

1. A process for preparing a sealed vial or cartridge containing a liquid pharmaceutical formulation ready for parenteral administration, comprising:
 - a. combining human parathyroid hormone and a buffer to maintain a pH from greater than 3 to 7, thereby forming a solution, and
 - b. sealing said solution in the vial or cartridge from which a therapeutically effective dose of parathyroid hormone can be withdrawn for use by a patient.
2. A pharmaceutical composition in the form of a solution, which comprises human parathyroid hormone and a buffer to maintain a pH from greater than 3 to 7, said solution being ready for parenteral administration in a human patient without having been freeze dried.
3. The pharmaceutical of claim 2, wherein said buffer is selected from citrate, tartrate, or acetate.
4. The pharmaceutical of claim 2, said solution further comprising a stabilizer.
5. The pharmaceutical of claim 4 wherein said stabilizer is mannitol.
6. The pharmaceutical of claim 2, wherein said human parathyroid hormone is selected from the group consisting of PTH (1-34), PTH (1-37), PTH (1-38), PTH (1-41) and PTH (1-84).
7. The pharmaceutical of claim 6, wherein said human parathyroid hormone is PTH (1-34).

8. A sealed vial or cartridge containing a pharmaceutical composition in the form of a solution ready for parenteral administration in a human patient, said formulation comprising:
 - a. human parathyroid hormone, and
 - b. a buffer to maintain a pH from greater than 3 to 7.
9. The vial or cartridge of claim 8 wherein said buffer is selected from citrate, tartrate, or acetate.
10. The vial or cartridge of claim 8, said formulation further comprising a stabilizer.
11. The vial or cartridge of claim 10 wherein said stabilizer is mannitol.
12. The vial or cartridge of claim 8, wherein said human parathyroid hormone is selected from the group consisting of PTH (1-34), PTH (1-37), PTH (1-38), PTH (1-41) and PTH (1-84).
13. The vial or cartridge of claim 12, wherein said human parathyroid hormone is PTH (1-34).
14. The vial or cartridge of claim 8 wherein said parathyroid hormone is at a concentration of 25 µg/ml to 1000 µg/ml.

SUMMARY OF THE INVENTION

According to one aspect of the present invention, there is provided a process for preparing a sealed vial or cartridge containing a liquid pharmaceutical formulation ready for parenteral administration, comprising combining human parathyroid hormone and a buffer to maintain a pH from greater than 3 to 7, thereby forming a solution, and sealing said solution in the vial or cartridge from which a therapeutically effective dose of parathyroid hormone can be withdrawn for use by a patient.

According to again another aspect of the present invention, there is provide a pharmaceutical composition in the form of a solution, which comprises human parathyroid hormone and a buffer to maintain a pH from greater than 3 to 7, said solution being ready for parenteral administration in a human patient without having been freeze dried.

According to yet another aspect of the present invention, there is provided a sealed vial or cartridge containing a pharmaceutical composition in the form of a solution ready for parenteral administration in a human patient, said formulation comprising human parathyroid hormone, and a buffer to maintain a pH from greater than 3 to 7.

According to yet another aspect of the present invention, there is provided a sealed vial or cartridge containing a pharmaceutical composition in the form of a solution comprising human parathyroid hormone, wherein the parathyroid hormone is selected from the group consisting of PTH (1-34), PTH (1-37), PTH (1-38), PTH (1-41) and PTH (1-84).

DETAILED DESCRIPTION

The invention relates to parathyroid hormone solutions that exhibit storage stability in terms of hormone composition and activity.

As active ingredient, the composition or solution may incorporate the full length, 84 amino acid form of parathyroid hormone, particularly the human form, hPTH (1-84) (SEQ ID NO: 1), obtained either recombinantly, by peptide synthesis or by extraction from human fluid. See, for example, U.S. Pat. No. 5,208,041, incorporated herein by reference. The amino acid sequence for hPTH (1-84) is reported by Kimura et al. in *Biochem. Biophys. Res. Comm.*, 114(2):493 (SEQ ID NO: 1).

The composition or solution may also incorporate as active ingredient fragments or variants of fragments of human PTH or of rat, porcine or bovine PTH that have human PTH activity as determined in the ovariectomized rat model of osteoporosis reported by Kimmel et al., *Endocrinology*, 1993, 32(4):1577.

The parathyroid hormone fragments desirably incorporate at least the first 34 N-terminal residues, such as PTH(1-34) (SEQ ID NO: 2), PTH(1-37), PTH(1-38) and PTH(1-41). Alternatives in the form of PTH variants incorporate from 1 to 5 amino acid substitutions that improve PTH stability and half-life, such as the replacement of methionine residues at positions 8 and/or 18 with leucine or other hydrophobic amino acid that improves PTH stability against oxidation and the replacement of amino acids in the 25-27 region with trypsin-insensitive amino acids such as histidine or other amino acid that improves PTH stability against protease. These forms of PTH are embraced by the term "parathyroid hormone" as used generically herein. The preferred hormone is human PTH(1-34) (SEQ ID NO: 2) also known as teriparatide. The hormones may be obtained by known recombinant or synthetic methods, such as described in U.S. Pat. No. 4,086,196, incorporated herein by reference.

The stabilizing agent incorporated into the solution or composition includes a polyol which includes a saccharide, preferably a monosaccharide or disaccharide, e.g., glucose, trehalose, raffinose, or sucrose; a sugar alcohol such as, for example,

mannitol, sorbitol or inositol, and a polyhydric alcohol such as glycerine or propylene glycol or mixtures thereof. A preferred polyol is mannitol or propylene glycol. The concentration of polyol may range from about 1 to about 20 wt-%, preferably about 3 to 10 wt-% of the total solution.

The buffering agent employed in the solution or composition of the present invention may be any acid or salt combination which is pharmaceutically acceptable and capable of maintaining the aqueous solution at a pH range of 3 to 7, preferably 3-6. Useful buffering systems are, for example, acetate, tartrate or citrate sources. Preferred buffer systems are acetate or tartrate sources, most preferred is an acetate source. The concentration of buffer may be in the range of about 2 mM to about 500 mM, preferably about 2 mM to 100 mM.

The stabilized solution or composition of the present invention may also include a parenterally acceptable preservative. Such preservatives include, for example, cresols, benzyl alcohol, phenol, benzalkonium chloride, benzethonium chloride, chlorobutanol, phenylethyl alcohol, methyl paraben, propyl paraben, thimerosal and phenylmercuric nitrate and acetate. A preferred preservative is m-cresol or benzyl alcohol; most preferred is m-cresol. The amount of preservative employed may range from about 0.1 to about 2 wt-%, preferably about 0.3 to about 1.0 wt-% of the total solution.

Thus, the present invention has provided, for example, a stabilized teriparatide solution containing mannitol, acetate and m-cresol with a predicted shelf-life of over 15 months at 5°C.

The parathyroid hormone compositions of the present invention may, if desired, be provided in a powder form containing not more than 2% water by weight, that results from the freeze-drying of a sterile, aqueous hormone solution prepared by mixing the selected parathyroid hormone, a buffering agent and a stabilizing agent as above described. Especially useful as a buffering agent when preparing lyophilized powders is a tartrate source. Particularly useful stabilizing agents include glycine, mannitol, sucrose, trehalose, raffinose or a mixture thereof.

The PTH solution and composition of the present invention incorporate PTH in a medically effective amount, a term used with reference to amounts useful either

therapeutically or in medical diagnosis. The particular amount of parathyroid hormone incorporated in the preparation can be pre-determined based on the type of PTH selected and on the intended end-use of the preparation. In one application, the preparations are exploited for therapeutic purposes, and particularly for the treatment of osteoporosis. Osteoporosis therapy entails administration of the reconstituted preparation by injection, desirably subcutaneous injection, in unit doses that reflect the prescribed treatment regimen but are, by way of example, for human PTH(1-34) (SEQ ID NO: 2), within the range from 25 µg PTH/mL of injected solution to 1000 µg/mL of injected solution per patient, with injection volumes being desirably from 0.02 to 1.3 mL. Accordingly, the purified PTH is desirably incorporated with the buffering agent and excipient to form an aqueous solution containing PTH in a concentration range from 25 µg/mL to 1000 µg/mL, preferably 100 µg/mL to 500 µg/mL, which is then sterile-filtered and filled into a vial or cartridge for use.

Once the preparation is obtained as an aqueous solution containing desired amounts and concentrations of the buffering agent, excipient and PTH, individual vials are filled with the solution to the desired volume. The advantage of the present invention is that the above solution may be prepared with sterile water without the need to undergo a freeze-drying process.

In an alternative embodiment of the invention, the preparations are provided in a form that yields a unit container of 100-500 µg human PTH(1-34) (SEQ ID NO: 2) upon reconstitution into about 1 mL (0.8-1.2 mL) of the reconstitution vehicle, and the vials are accordingly loaded with about 1 mL of the aqueous PTH preparation, for subsequent freeze-drying.

In a preferred alternative embodiment of the invention, the PTH preparation subjected to freeze-drying comprises from 25 to 1000 µg/mL of human PTH(1-34) (SEQ ID NO: 2), from 2 to 8% by weight of mannitol, and a tartrate source in an amount capable of buffering the preparation to within the range from 3.0 to 6.5 upon reconstitution in sterile water. In specific embodiments of the invention, the tartrate buffering agent is incorporated in an amount sufficient to buffer the pH to 3.5 to 5.5.

In addition to their therapeutic use, the present PTH composition can be formulated and administered to aid in medical diagnosis, and particularly to assist in

establishing the diagnosis of hypoparathyroidism and pseudohypoparathyroidism in hypocalcemic patients. Except for the dose of PTH, the composition of the PTH preparation will remain as described herein for therapeutic use. An intravenously infused, single dose of human PTH(1-34) (SEQ ID NO: 2) that is equal to 200 International Units of PTH activity is appropriate for this diagnostic purpose. Diagnosis is then made by determining the effect of administered PTH or urinary cAMP levels, with cAMP elevation being indicative of the hypoparathyroidism condition, rather than its pseudoform.

The examples which follow are illustrative of the invention and are not intended to be limiting.

EXAMPLES

Example 1

0.1 mg rhPTH (1-34) (SEQ ID NO: 2), 50 mg mannitol, 2.5 mg m-cresol, 0.52 mg acetic acid and 0.12 mg sodium acetate were mixed into a solution with 1 ml of distilled water.

Example 2

0.25 mg rhPTH (1-34) (SEQ ID NO: 2), 45.4 mg mannitol, 3 mg m-cresol, 0.41 mg acetic acid and 0.1 mg sodium acetate were mixed into a solution with 1 ml of distilled water.

The formulations of the present invention, Examples 1 and 2 were compared to solutions containing no stabilizer, 0.9% NaCl, 20 mM acetate and 10 mM acetate as primary stabilizer. The stability was measured by determining the amount in % of rhPTH (1-34) (SEQ ID NO: 2) remaining after a certain time. The measurement was made by HPLC. The results are shown in Tables 1 and 2.

Table 1

Effect of Primary Stabilizer on Chemical Stability
of rhPTH (1-34) at 50°C

	Water	0.9% NaCl	20 mM acetate	10 mM acetate
Time, days	% Remaining			
Initial	100	100	100	100
7	74	81	84	80
14	55	58	67	71

Table 2

Comparison of Stability of rhPTH (1-34) at 30°C

	20 mM acetate	10 mM acetate	Example 1	Example 2
Time, days	% Remaining			
Initial	100	100	100	100
7	96	94	100	--
14	94	92	96	100
21	90	93	97	--
30	--	81	96	96

Example 3

The following experiment was carried out to show that lyophilized powder formulations prepared from stabilized solutions of the present invention are more stable than a control which was prepared from PTH(1-34) and mannitol alone.

A control solution and solutions for samples A through O were prepared as previously described with the ingredients and concentrations shown in Table 3. The solutions were then freeze-dried and the resulting lyophilized powder formulations were stored at 40°C for a one month period. The amount of PTH(1-34) remaining in each sample was then measured by HPLC. The results are shown in Table 3.

Table 3Stability of PTH(1-34) Lyophilized Formulations at 40°C for One Month

Sample	PTH(1-34) mg/mL	Bulking Agent	Bulking Agent Conc. mg/mL	Buffer	Buffer Conc. mM	% PTH Remaining
Control	0.2	mannitol	40	--	--	78
A	0.5	mannitol	30	acetate	5	90
B	0.5	glycine	30	acetate	5	98
C	0.5	sucrose	30	acetate	5	98
D	0.5	trehalose	30	acetate	5	97
E	0.5	raffinose	30	acetate	5	99
F	0.75	mannitol	30	tartrate	15	95
G	1.5	sucrose & mannitol	5/25	tartrate	5	99
H	0.75	sucrose & mannitol	5/25	tartrate	15	99
I	1.5	mannitol	30	tartrate	5	96
J	1.5	sucrose	30	tartrate	15	100
K	1.5	mannitol	30	tartrate	15	99
L	0.75	sucrose	30	tartrate	15	100
M	0.75	sucrose	30	tartrate	5	100
N	1.5	sucrose & mannitol	5/25	tartrate	15	99
O	1.5	sucrose & mannitol	5/25	acetate	5	91*

* the stability at 2 months was 96%