A novel form of ibandronate sodium which is particularly suitable for pharmaceutical applications, and a process for preparing said novel form.
IBANDRONATE SODIUM PROPYLENE GLYCOL SOLVATE AND PROCESSES FOR THE PREPARATION THEREOF

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This is a divisional application of U.S. application Ser. No. 11/639,204, filed on Dec. 15, 2006.

FIELD OF THE INVENTION

[0002] The present invention relates to a new solvated form of ibandronate sodium (Boniva®) by formation of a 1:1 ibandronate sodium propylene glycol solvate, and a method for its preparation. This form is particularly well-suited for pharmaceutical applications.

BACKGROUND OF THE INVENTION

[0003] Ibandronate sodium (1,3-(N-methyl-N-pentyl) amino-1-hydroxypropane-1,1-diphosphonic acid, monosodium salt) is a third-generation member of the bis-phosphonate class of drugs effective for the treatment of bone disorders such as osteoporosis. It is marketed under the brand name Boniva and its mode of action is to inhibit osteoclast-mediated bone resorption. An advantage of this pharmaceutical is that it can be used as a once-monthly treatment.
Figure 1, Ibandronate Sodium (Boniva®)
The use of ibandronate sodium as a medicine was originally disclosed in U.S. Pat. No. 4,927,814 assigned to Boehringer Mannheim GmbH. This patent teaches the use of various structurally related diphasonic acid derivatives, including ibandronate, for the treatment of various bone disorders. This patent also teaches the physiologically active salts of these diphasonic acids, in particular their mono- or dialkali metal salts and states that these alkali metal salts can usually be readily purified by recrystallization from water/methanol or from water/acetic acid.

WO 2005/063779 (Lyogen) discloses improved processes to prepare 10 diphasonic acids, in particular Risendronate, Zoledronate and Ibandronate acids. It also discloses an amorphous form of ibandronate sodium. In contrast to the prior art, this patent application teaches the use of phosphorus oxychloride and phosphoric acid in a well-defined ratio to obtain a fluid mass that was stirrable throughout the reaction. Similarly, U.S. Pat. No. 5,908,959 (Apotex Inc.) resolved stirring issues related to another diphasonic acid, Alendronate and its salts, by performing the phosphorylation reaction in the presence of polyethylene glycol.

US patent application 2006/0172975 (Hoffmann-La Roche) describes ibandronate sodium monohydrate Form A and Form B.

WO 2006/024024 (Teva) teaches solid amorphous and crystalline forms of ibandronate sodium which are often full or partial solvates and hydrates. Various primary alcohols are listed as possible solvating agents including ethanol and butanol. Of note is that many of the types of solvating agents are toxic at higher doses and therefore would have to be removed before they could be used as an active ingredient. To this end, there are well-recognized international guidelines regarding the amount of residual solvent permissible in active ingredients [ICH Q3C(R3) in http://www.ich.org/LOB/media/MEDIA4423.pdf].

Teva application WO 2006/024024 also teaches that common liquid carriers such as propylene glycol can be used to suspend or dissolve their active forms of ibandronate sodium and solid excipients for liquid pharmaceutical compositions. This application, however, does not contemplate the novel propylene glycol solvate form of the present invention.

The use of propylene glycol as a solvating agent for certain pharmaceuticals is known, for instance Celecoxib (US 2006/0052432), Olanzapine (US 2006/0233794), Eplerenone (US 2005/0267302), Azithromycin (U.S. Pat. No. 7,105,179) and Cefepime derivatives (U.S. Pat. No. 4,091,213). However, the use of propylene glycol for the diphasonic class of drugs, including ibandronate sodium, is not described in these prior art references.

Another patent application by Teva, WO 2006/002348, discloses various routes to ibandronate acid and various crystalline and amorphous forms of ibandronate acid. This application also teaches the benefits of propylene glycol as a carrier for liquid pharmaceutical compositions of ibandronate acid, but does not contemplate the novel ibandronate sodium propylene glycol solvate of the present invention.

There are various patents which disclose formulation improvements in order to minimize well-known tolerability problems associated with the diphasonic class of drugs. For instance, U.S. Pat. No. 6,143,326 (Roche Diagnostics) and U.S. Pat. No. 6,294,196 (Hoffmann-La Roche), relate to the outer coat of the tablets containing ibandronate sodium in order to obtain the desired release profile.

Another patent is U.S. Pat. No. 6,468,559 (Lipocine) which describes an effective oral dosage form of many bisphosphonates (14), including ibandronate acid and salts, using enterically coated capsules comprised of the active and a pharmaceutically acceptable, substantially non-aqueous liquid or semi-solid carrier in which the active agent is dissolved or suspended. There are numerous examples (>200) of the non-aqueous carriers mentioned in U.S. Pat. No. 6,468,559, including propylene glycol. In the teachings of this patent, the active is dissolved or suspended in the liquid carriers, but is not made into, or isolated as, a solvate.

Given the difficulties associated with finding suitable processes to and forms of ibandronate sodium, new and industry acceptable solutions, which offer advantages relative to the prior art, were required.

**SUMMARY OF THE INVENTION**

During our process optimization work to find novel, cost-effective and robust synthetic procedures to ibandronate sodium and improved forms of the active ingredient, we surprisingly discovered that a diol, namely propylene glycol (1,2-propanediol), forms a crystalline 1:1 solvate with ibandronate sodium that can be easily purified and, thereafter, formulated into effective dosage forms.

The use of propylene glycol as a solvating agent has many advantages. These include that it is inexpensive and widely-available, having many industrial applications including use as a moisturizer for medicines, cosmetics, food and tobacco products and as a humectant food additive. Furthermore, it has an established safety and can be purchased as pharmaceutically acceptable NF-grade material. Moreover, its use is listed as a Food Additive in FDA’s “Everything Added to Food in the United States (EAFUS)” database (http://www.cfsan.fda.gov/~dms/eafus.html).

The ibandronate sodium propylene glycol solvate of this invention is easily made and crystalline. This form has many desirable characteristics including that it is non-hygroscopic, free-flowing, and is chemically and polymorphically stable. Another benefit is that it is easily dried, even on industrial scale, since it retains very little residual solvent.

Propylene glycol can be obtained in either enantiomer enriched R or S form or as a mixture of enantiomers (i.e., as a racemate). The latter racemate form of propylene glycol is more preferred for this invention.

In another aspect of the invention, the phosphorylation of 3-(N-methyl-N-pentylamino)propanoic acid (or its hydrochloride salt) using phosphoric acid and phosphorous trichloride in the presence of polyethylene glycol led to superior results. Most significantly, it allowed for the reaction to be accomplished with effective stirring throughout thereby permitting facile and safe scale-up to industrial levels.

**BRIEF DESCRIPTION OF THE DRAWINGS**

**Fig. 1** illustrates a powder x-ray diffraction pattern of ibandronate sodium propylene glycolate.

**Fig. 2** illustrates a DSC thermogram of crystalline ibandronate sodium propylene glycolate.

**Fig. 3** illustrates an IR spectrum of ibandronate sodium propylene glycolate.

**DETAILED DESCRIPTION OF THE INVENTION**

The solvated form of ibandronate sodium may be formed in various ways. For instance, ibandronate sodium...
can be dissolved in water at about 30°C to about 100°C, more preferably at about 50°C to about 80°C. Typically this requires about 3 volumes of water relative to the weight of the Ibandronate sodium. This is followed by the addition of about 0.5 to 2 volumes of propylene glycol and the solution is cooled at about −5°C to about 30°C, more preferably at about 0°C to about 20°C, most preferably at about 0°C to about 5°C. The material is isolated by filtration and rinsed with an organic solvent selected from a C3 to C5 ester, most preferably ethyl acetate or a C3 to C6 alkyl ketone, most preferably aceton, or a C4 to C8 cyclic or acyclic ether, most preferably tetrahydrofuran. This last step is done to remove the excess propylene glycol.

The Ibandronate sodium propylene glycolate prepared by this method can be characterized by a PXRD pattern having characteristic peaks expressed in angle 2-theta at approximately:

<table>
<thead>
<tr>
<th>Angle 2-theta</th>
<th>Intensity %</th>
</tr>
</thead>
<tbody>
<tr>
<td>9.7</td>
<td>0.24</td>
</tr>
<tr>
<td>11.8</td>
<td>0.49</td>
</tr>
<tr>
<td>18.3</td>
<td>0.86</td>
</tr>
<tr>
<td>19.4</td>
<td>0.57</td>
</tr>
<tr>
<td>19.8</td>
<td>0.83</td>
</tr>
<tr>
<td>24.5</td>
<td>0.59</td>
</tr>
</tbody>
</table>

The Ibandronate sodium propylene glycolate prepared by this method can be further characterized by a PXRD pattern as shown in FIG. 1.

The Ibandronate sodium propylene glycolate prepared by this method can be characterized by a DSC having a major endotherm at a peak onset temperature of about 220°C and a peak maximum of about 228°C. The crystalline Ibandronate sodium propylene glycolate prepared by this method can be further characterized by a DSC thermogram as shown in FIG. 2.

The Ibandronate sodium propylene glycolate prepared by this method can be characterized by its IR spectrum (1% KBr) having characteristic peaks expressed in cm$^{-1}$ at approximately:

<table>
<thead>
<tr>
<th>IR band (cm$^{-1}$)</th>
<th>% Transmission</th>
</tr>
</thead>
<tbody>
<tr>
<td>3382</td>
<td>31</td>
</tr>
<tr>
<td>2385</td>
<td>34</td>
</tr>
<tr>
<td>1467</td>
<td>42</td>
</tr>
<tr>
<td>1380</td>
<td>46</td>
</tr>
<tr>
<td>1162</td>
<td>3.2</td>
</tr>
<tr>
<td>1032</td>
<td>4.8</td>
</tr>
<tr>
<td>931</td>
<td>20</td>
</tr>
</tbody>
</table>

The Ibandronate sodium propylene glycolate prepared by this method can be further characterized by an IR spectrum (taken in 1% KBr) as depicted in FIG. 3.

In another embodiment of the invention, the Ibandronate sodium is directly isolated by addition of propylene glycol to a mixture of Ibandronic acid in a substantially aqueous media, adjusting the pH to about 4 to about 5 and then adding about 0.5 to about 2 volumes propylene glycol. The solution is cooled at about −5°C to about 30°C, more preferably at about 0°C to about 20°C, most preferably at about 0°C to about 5°C. The material is isolated by filtration and rinsed with an organic solvent selected from a C3 to C5 ester, most preferably ethyl acetate or a C3 to C6 alkyl ketone, most preferably aceton, or a C4 to C8 cyclic or acyclic ether, most preferably tetrahydrofuran. This last step is done to remove the excess propylene glycol.

In another embodiment of the invention, an improved procedure for the phosphorylation of 3-(N-methyl-N-pentylamino)propanoic acid (or its hydrochloride salt) was accomplished using phosphorus acid and phosphorous trichloride in the presence of polyethylene glycol media. This permitted the reaction to be accomplished with effective stirring throughout and was easily scaled, in a safe manner, to multi-kilo levels. The most performed average molecular weight of polyethylene glycol was about 400 g/mol. The amount of phosphorus acid and phosphorous trichloride relative to the 3-(N-methyl-N-pentylamino)propanoic acid substrate was about 0.8 to about 2.0 equivalents, more preferably about 1.0 to about 1.6 equivalents. The preferred temperature for performing the reaction was about 45 to about 70°C, more preferably at about 55 to about 60°C. The most preferably amount of polyethylene glycol was about 0.8 to about 2 volumes, relative to the weight of the 3-(N-methyl-N-pentylamino)propanoic acid substrate. This reaction is also preferably performed in the presence of a co-solvent, most preferably toluene. The preferred amount is about 2 to about 4 volumes, relative to the weight of the 3-(N-methyl-N-pentylamino)propanoic acid substrate.

The following examples are representative of the present invention and are not intended to be limiting.

**EXAMPLE 1**

Preparation of Sodium Ibandronate Propylene Glycolate from 3-(N-Methyl-N-Pentylamino)Propanoic Acid Hydrochloride

3-(N-Methyl-N-pentylamino)propanoic acid hydrochloride (100 g) was suspended in toluene (300 mL) and polyethylene glycol 400 (120 mL). Phosphorous acid (43.01 g 1.1 eq.) was added to the mixture. The mixture was warmed and phosphorous trichloride (98.22 g, 1.5 eq.) was added at a rate such that the temperature remained below 60°C. The mixture was stirred for 10 hours at 55-60°C whereupon it was quenched by adding water (450 mL). The water addition was controlled to maintain a reaction temperature below 70°C. The layers were separated and the aqueous layer was refluxed for 6 hours. The mixture was cooled to 40-45°C to provide Ibandronic acid in solution and the pH was adjusted to 4.3-4.5 using 50% aqueous sodium hydroxide. Propylene glycol (150 mL) was then added to the reaction mixture at 60-65°C and stirred for 4-5 hours. The mixture was cooled to 20-25°C over a period of 2-3 hours and then for 3-4 hours at 0-5°C, filtered, washed with acetone (2×150 mL) and dried in vacuo.

Sodium Ibandronate (145 g) from above was suspended in 363 mL (2.6 vol) of water in a round bottom flask. The flask was heated to dissolution (60-65°C) whereupon propylene glycol (72 mL) was added to the flask and held at this temperature for 6 hours. The flask was cooled to 20-25°C and then 0-5°C and held at this temperature for 3 hours. The precipitated solid was isolated by filtration and rinsed with ethyl acetate (2×200 mL). The damp filter cake was then stirred with ethyl acetate (650 mL) for 3 hours at 40-45°C...
C., isolated by filtration and dried in a vacuum oven at 60-65° C. This provided 110.3 grams of Ibandronate sodium propylene glycolate.

0033  Sodium Ibandronate propylene glycolate having the following analytical characteristics was obtained.

0034  1H-NMR (400 MHz, D$_2$O): δ=3.90-3.82 (1H, m); 3.59-3.50 (2H, m), 3.45-3.40 (1H, m), 3.36-3.30 (1H, m), 3.27-3.18 (1H, m), 3.07-3.01 (1H, m), 2.84 (3H, s), 2.44-2.26 (2H, m), 1.74-1.68 (2H, m), 1.35-1.29 (4H, m), 1.13 (3H, d, J=6.7 Hz), 0.88 (3H, t, J=6.9 Hz).

0035  13C-NMR (100 MHz, D$_2$O): δ=75.0 (t, J=135.4 Hz), 70.6, 69.3, 59.0, 55.6 (t, J=6.8 Hz), 42.1, 30.64, 30.55, 25.9, 24.2, 20.8, 15.8.

0036  Mass Spectroscopy (m/z, ES$^-$): 318 (M-Na, 100).

0037  Elemental Analysis: Calculated for C$_{12}$H$_{16}$O$_4$N$_2$: C 34.54; H 7.25; N 3.36. Found: C 34.44; H 7.56; N 3.33.

0038  IR (1% KBr): 3382, 3160, 2962, 2863, 2385, 1482, 1468, 1381, 1344, 1210, 1162, 1090, 1063, 1032 cm$^{-1}$.

EXAMPLE 2
Preparation of Sodium Ibandronate Propylene Glycolate from Ibandron Acid

0039  Ibandron acid (6.30 g, 19.7 mmol) was suspended in water (18.0 mL) and heated to dissolution (42° C.). The pH was adjusted to 4.4 using 50% aqueous sodium hydroxide. 1,2-Propanediol was added to the reaction mixture at 56° C. and the mixture was stirred for 2.5 hours. The mixture was cooled to 20-25° C. over a period of 2.5 hours and stirred for another 15 hours. The mixture was filtered, washed with ethyl acetate (2×10 mL). The damp solid was suspended in 30 mL of ethyl acetate and stirred at 20-25 °C. for 1 hour, filtered, rinsed with ethyl acetate (2×10 mL) and dried in vacuo at 65° C. for 8 hours to yield sodium Ibandronate propylene glycolate (4.54 g).

EXAMPLE 3
Preparation of Sodium Ibandronate R-Propylene Glycolate by Recrystallization

0040  Sodium Ibandronate (2.0 g) was suspended in 6 mL (3.0 vol) of water in a round bottom flask. The flask was heated to dissolution (55-65° C) at which point 2 mL R-propylene glycol (1.0 vol) was added to the flask. The temperature of the flask was held at 60-65° C for 2 hours, then cooled to 20-25° C. and held at this temperature for 0.5 hours. Acetone (4 mL, 2.0 vol) was added and the flask was stirred at 20-25° C. for 0.5 hours. The precipitated solid was isolated by filtration, rinsed with acetone and dried in a vacuum oven at 60-65° C. This provided 1.97 grams of Ibandronate sodium R-propylene glycolate.

EXAMPLE 4
Preparation of Sodium Ibandronate S-Propylene Glycolate by Recrystallization

0041  Sodium Ibandronate (2.0 g) was suspended in 6 mL (3.0 vol) of water in a round bottom flask. The flask was heated to dissolution (55-65° C) at which point 2 mL S-propylene glycol (1.0 vol) was added to the flask. The temperature of the flask was held at 60-65° C for 2 hours, then cooled to 20-25° C. and held at this temperature for 0.5 hours. Acetone (4 mL, 2.0 vol) was added and the flask was stirred at 20-25° C. for 0.5 hours. The precipitated solid was isolated by filtration, rinsed with acetone and dried in a vacuum oven at 60-65° C. This provided 2.05 grams of Ibandronate sodium S-propylene glycolate.

0042  As many changes can be made to the examples which exemplify the invention without departing from the scope of the invention, it is intended that all matter contained herein be considered illustrative of the invention and not in a limiting sense.

1. The preparation of Ibandron acid and its sodium salt comprising reacting 3-(N-methyl-pentylamino)propanoic acid or its hydrochloride salt, phosphorous acid and phosphorous trichloride in the presence of polyethylene glycol.

2. The preparation of Ibandron acid and its sodium salt according to claim 1 wherein the polyethylene glycol has an average molecular weight of about 400.

3. The preparation of Ibandron acid and its sodium salt according to claim 1 or 2 wherein the phosphorylation reaction is performed at about 45° C. to about 70° C.

4. The preparation of Ibandron acid and its sodium salt according to claim 1 or 2 wherein the phosphorylation reaction is performed at about 55° C. to about 60° C.

5. The preparation of Ibandron acid and its sodium salt according to claim 3 or 4 wherein the phosphorylation reaction is performed in about 0.8 to about 2 volumes of polyethylene glycol.

6. The preparation of Ibandron acid and its sodium salt according to claim 3 or 4 wherein the phosphorylation reaction is performed using about 0.8 to about 2 equivalents of phosphorous acid and phosphorous trichloride relative to the amount of 3-(N-methyl-N-pentylamino)propanoic acid substrate.

7. The preparation of Ibandron acid and its sodium salt according to claim 3 or 4 wherein the phosphorylation reaction is performed using about 1.0 to about 1.6 equivalents of phosphorous acid and phosphorous trichloride relative to the amount of 3-(N-methyl-N-pentylamino)propanoic acid substrate.

8. The preparation of Ibandron acid and its sodium salt according to claims 1, 2 or 3 wherein the phosphorylation reaction is performed in the presence of a co-solvent.

9. The preparation of Ibandron acid and its sodium salt according to claim 8 wherein the phosphorylation reaction is performed in the presence of toluene as a co-solvent.

10. The preparation of Ibandron acid and its sodium salt according to claim 9 wherein the phosphorylation reaction is performed in a toluene co-solvent where the amount of toluene is about 2 to about 4 volumes, relative to the weight of the 3-(N-methyl-N-pentylamino)propanoic acid substrate.

11. The preparation of Ibandronate sodium according to claims 1, 2 or 8 wherein the Ibandronate sodium is further converted to Ibandronate sodium propylene glycolate.

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