



NDA 21976/S-022

**SUPPLEMENT APPROVAL**

Janssen Products, L.P.  
Attention: Charles Zezza, Ph.D.  
Director, Global Regulatory Affairs  
920 Route 202 S  
Raritan, NJ 08869

Dear Dr. Zezza:

Please refer to your Supplemental New Drug Application (sNDA) dated and received November 17, 2011, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for Prezista<sup>®</sup> (darunavir) 75 mg, 150 mg, 400 mg, and 600 mg Tablets

We acknowledge receipt of your amendments dated January 6, and February 7, 2012 in response to our General Advice letter dated, December 12, 2011 and our February 3, 2012 e-mail comments.

We also refer to our letter dated October 19, 2011, notifying you, under Section 505(o)(4) of the FDCA, of new safety information that we believe should be included in the labeling for antiretroviral products. This information pertains to the risk of the autoimmune disorder as syndromes that can occur in the setting of immune reconstitution with the use of antiretroviral products.

In addition, we refer to non-safety labeling changes in our October 19, 2011 letter for all antiretroviral products based on recent studies demonstrating decreased transmission of HIV when HIV-infected patients or their uninfected partners take antiretroviral medication.

This supplemental new drug application provides for revisions to the labeling for Prezista<sup>®</sup> (darunavir) 75 mg, 150 mg, 400 mg, and 600 mg Tablets, consistent with our October 19 and December 12, 2011 letters and February 3, 2012 e-mail comments, as follows (additions are noted by underline and deletions are noted by ~~strike through~~).

1. The **RECENT MAJOR CHANGES** in the Highlights section of the labeling has been revised as follows:

-----**RECENT MAJOR CHANGES**-----

- Indications and Usage
  - Pediatric Patients (1.2)      12/2011
- Dosage and Administration

- Adult Patients (2.1) 12/2011
  - Pediatric Patients (2.2) 12/2011
  - Warnings and Precautions
    - Severe Skin Reactions (5.3) 10/2011
    - Immune Reconstitution Syndrome (5.8) 02/2012
    - Pediatric Patients (5.11) 12/2011
2. The revision date has been changed from 12/2011 to 02/2012 throughout the label.
3. The **WARNINGS AND PRECAUTIONS/Immune Reconstitution Syndrome** sub-section has been revised as follows:

Immune reconstitution syndrome has been reported in patients treated with combination antiretroviral therapy, including PREZISTA. During the initial phase of combination antiretroviral treatment, patients whose immune systems responding to antiretroviral therapy may develop an inflammatory response to indolent or residual opportunistic infections (such as Mycobacterium avium infection complex, cytomegalovirus, Pneumocystis jirovecii pneumonia (PCP), and/or tuberculosis), which may necessitate further evaluation and treatment.

Autoimmune disorders (such as Graves' disease, polymyositis, and Guillain-Barré syndrome) have also been reported to occur in the setting of immune reconstitution; however, the time to onset is more variable, and can occur many months after initiation of treatment.

4. The Table 9 in the **7.3 Established and Other Potentially Significant Drug Interactions** section of the package insert has been revised as follows:

| <b>Table 9: Established and Other Potentially Significant Drug Interactions:<br/>Alterations in Dose or Regimen May Be Recommended<br/>Based on Drug Interaction Studies or Predicted Interaction</b><br>[See <i>Clinical Pharmacology</i> (12.3) for Magnitude of Interaction, Tables 12 and 13] |   |  |
|---|---|--|
| <b>Concomitant Drug Class:<br/>Drug Name</b>  | <b>Effect on Concentration of<br/>Lopinavir or Concomitant<br/>Drug</b> | <b>Clinical Comment</b>  |
| <i>Other Agents</i>   |   |  |
| <b><u>HMG-CoA Reductase<br/>Inhibitors:</u></b><br><u>pravastatin</u><br><u>atorvastatin</u><br><u>rosuvastatin</u>   | ↑ pravastatin<br>↑ atorvastatin<br>↑ rosuvastatin                       | <u>Titrate atorvastatin, pravastatin or rosuvastatin dose carefully and use the lowest possible necessary dose while monitoring for safety. Caution should be used when the Do not exceed atorvastatin dose exceeds 20 mg/day of atorvastatin, pravastatin or rosuvastatin with careful monitoring, or consider other HMG CoA reductase inhibitors such as fluvastatin in combination with PREZISTA/ritonavir.</u> |

5. The **PATIENT COUNSELING INFORMATION/Information About Therapy with PREZISTA** sub-section has been revised as follows:

~~Patients should be informed that PREZISTA is not a cure for HIV infection. Patients should stay on continuous HIV therapy to control HIV infection and decrease HIV-related illnesses.~~

~~Patients should be told that sustained decreases in plasma HIV RNA have been associated with a reduced risk of progression to AIDS and death. Patients should remain under the care of a physician while using PREZISTA. Patients should be advised to continue to practice safer sex and to use latex or polyurethane condoms to lower the chance of sexual contact with any body fluids such as semen, vaginal secretions or blood. Patients should be advised never to share personal items that can have blood or body fluids on them, like toothbrushes and razor blades. Patients should be advised never to re-use or share needles. PREZISTA is not a cure for HIV-1 infection and patients may continue to experience illnesses associated with HIV-1 infection, including opportunistic infections. Patients should remain under the care of a physician when using PREZISTA.~~

Patients should be advised to avoid doing things that can spread HIV-1 infection to others.

- **Do not share needles or other injection equipment.**
- **Do not share personal items that can have blood or body fluids on them, like toothbrushes and razor blades.**
- **Do not have any kind of sex without protection.** Always practice safe sex by using a latex or polyurethane condom to lower the chance of sexual contact with semen, vaginal secretions, or blood.
- **Do not breastfeed.** We do not know if PREZISTA can be passed to the baby through breast milk and whether it could harm the baby. Also, mothers with HIV-1 should not breastfeed because HIV-1 can be passed to the baby in the breast milk.

6. **Patient Information:**

- a. The “**Does PREZISTA cure HIV or AIDS?**” section has been revised as follows:

**Does PREZISTA cure HIV or AIDS? What is PREZISTA?**

~~**PREZISTA does not cure HIV infection or AIDS.** People taking PREZISTA may still develop infections or other conditions associated with HIV infection, including opportunistic infections (e.g., pneumonia and herpes virus infections).~~

~~Patients must stay on continuous HIV therapy to control infection and decrease HIV-related illnesses.~~

~~Avoid doing things that can spread HIV-1 infection.~~

- ~~▪ **Do not share needles or other injection equipment.**~~
- ~~▪ **Do not share personal items that can have blood or body fluids on them, like toothbrushes and razor blades.**~~
- ~~▪ **Do not have any kind of sex without protection.** Always practice safe sex by using a latex or polyurethane condom to lower the chance of sexual contact with any body fluids such as semen, vaginal secretions, or blood. Never re-use or share needles.~~

~~Ask your healthcare provider if you have any questions on how to prevent passing HIV to other people. **PREZISTA does not cure HIV infection or AIDS and you may continue to experience illnesses associated with HIV-1 infection, including opportunistic infections. You should remain under the care of a doctor when using PREZISTA.**~~

Avoid doing things that can spread HIV-1 infection.

- **Do not share needles or other injection equipment.**
- **Do not share personal items that can have blood or body fluids on them, like toothbrushes and razor blades.**
- **Do not have any kind of sex without protection.** Always practice safe sex by using a latex or polyurethane condom to lower the chance of sexual contact with semen, vaginal secretions, or blood.

Ask your healthcare provider if you have any questions on how to prevent passing HIV to other people.

- b. The “**Does PREZISTA reduce the risk of passing HIV to others?**” section has been deleted.
- c. The sixth bulleted paragraph in the “**What should I tell my doctor before taking PREZISTA?/ PREZISTA may not be right for you. Before taking PREZISTA, tell your doctor or healthcare provider if you:**” section has not been revised as follows:
  - ~~are breastfeeding or plan to breastfeed. Do not breastfeed if you are taking PREZISTA. You should not breastfeed if you have HIV because of the chance of passing HIV to your baby. Talk with your healthcare provider about the best way to feed your baby. The Centers for Disease Control and Prevention (CDC) recommends that HIV-infected mothers not breastfeed to avoid the risk of passing HIV infection to your baby. **Do not breastfeed. We do not know if PREZISTA can be passed to your baby in your breast milk and whether it could harm your baby. Also, mothers with HIV-1 should not breastfeed because HIV-1 can be passed to the baby in the breast milk.**~~

We have completed our review of this supplemental application, as amended. It is approved, effective on the date of this letter, for use as recommended in the enclosed, agreed-upon labeling text.

## **CONTENT OF LABELING**

As soon as possible, but no later than 14 days from the date of this letter, submit the content of labeling [21 CFR 314.50(l)] in structured product labeling (SPL) format using the FDA automated drug registration and listing system (eLIST), as described at <http://www.fda.gov/ForIndustry/DataStandards/StructuredProductLabeling/default.htm>. Content of labeling must be identical to the enclosed labeling (text for the package insert, text for the patient package insert), with the addition of any labeling changes in pending “Changes Being Effected” (CBE) supplements, as well as annual reportable changes not included in the enclosed labeling.

Information on submitting SPL files using eLIST may be found in the guidance for industry titled “SPL Standard for Content of Labeling Technical Qs and As” at <http://www.fda.gov/downloads/DrugsGuidanceComplianceRegulatoryInformation/Guidances/UCM072392.pdf>.

The SPL will be accessible from publicly available labeling repositories.

Also within 14 days, amend all pending supplemental applications for this NDA, including CBE supplements for which FDA has not yet issued an action letter, with the content of labeling [21 CFR 314.50(l)(1)(i)] in MS Word format, that includes the changes approved in this supplemental application, as well as annual reportable changes and annotate each change. To facilitate review of your submission, provide a highlighted or marked-up copy that shows all changes, as well as a clean Microsoft Word version. The marked-up copy should provide appropriate annotations, including supplement number(s) and annual report date(s).

## **PROMOTIONAL MATERIALS**

You may request advisory comments on proposed introductory advertising and promotional labeling. To do so, submit the following, in triplicate, (1) a cover letter requesting advisory comments, (2) the proposed materials in draft or mock-up form with annotated references, and (3) the package insert(s) to:

Food and Drug Administration  
Center for Drug Evaluation and Research  
Office of Prescription Drug Promotion (OPDP)  
5901-B Ammendale Road  
Beltsville, MD 20705-1266

You must submit final promotional materials and package insert(s), accompanied by a Form FDA 2253, at the time of initial dissemination or publication [21 CFR 314.81(b)(3)(i)]. Form FDA 2253 is available at <http://www.fda.gov/opacom/morechoices/fdaforms/cder.html>; instructions are provided on page 2 of the form. For more information about submission of promotional materials to the Office of Prescription Drug Promotion (OPDP), see <http://www.fda.gov/AboutFDA/CentersOffices/CDER/ucm090142.htm>.

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