



NDA 019386/S-043

**SUPPLEMENT APPROVAL**

Baxter Healthcare Corporation  
Attention: Amy Giertych  
Senior Director Global Regulatory Affairs  
1620 Waukegan Road, MPGR-AL  
McGaw Park, IL 60085

Dear Ms. Giertych:

Please refer to your Supplemental New Drug Application (sNDA) originally submitted December 30, 2010 and resubmitted February 17, 2012, under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for Brevibloc (esmolol hydrochloride), 10 mg per ml and 20 mg per ml for Injection.

The February 17, 2012 submission constituted a complete response to our July 1, 2011, action letter.

This "Prior Approval" supplemental new drug application provides for revisions to comply with the formatting requirements of 21 CFR 201.56(d). Content changes were also made in the following sections: 1, 2, 3, 4, 5, 6, 7, 8, 10, and 12. Other editorial and organizational changes were made throughout the label.

We have completed our review of this supplemental application. 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, 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).

### **REPORTING REQUIREMENTS**

We remind you that you must comply with reporting requirements for an approved NDA (21 CFR 314.80 and 314.81).

If you have any questions, please call Michael Monteleone, Regulatory Project Manager, at (301) 796-1952.

Sincerely,

*{See appended electronic signature page}*

Norman Stockbridge, MD, PhD  
Director  
Division of Cardiovascular and Renal Products  
Office of Drug Evaluation I  
Center for Drug Evaluation and Research

ENCLOSURE(S):

- Content of Labeling

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**This is a representation of an electronic record that was signed electronically and this page is the manifestation of the electronic signature.**  
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/s/  
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NORMAN L STOCKBRIDGE  
12/11/2012