



NDA 208610-S-006  
NDA 208611-S-005

## SUPPLEMENT APPROVAL

Melinta Therapeutics, Inc.  
Attention: Starr Shangle  
Senior Director, Regulatory Affairs  
300 Tri-State International, Suite 272  
Lincolnshire, IL 60069

Dear Ms. Shangle,

Please refer to your Supplemental New Drug Applications (sNDAs) dated and received January 18, 2019 and your amendments, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for the following;

NDA 208610 Baxdela (delafloxacin) 450 mg Tablets  
NDA 208611 Baxdela (delafloxacin) 300 mg Injection

We also refer to our letter dated December 20, 2018, notifying you, under Section 505(o)(4) of the FDCA, of new safety information that we believe should be included in the labeling for the systemic fluoroquinolone class of antibacterial drugs. This information pertains to the risk of aortic aneurysm and dissection.

We also refer to our February 13, 2019 letter that informed you of an additional extension of the discussion period to allow us to complete our review and reach agreement on the content of the labeling.

These supplemental new drug applications provide for the inclusion of new safety information describing the risk of aortic aneurysm and dissection in the following sections of the labeling.

- **WARNINGS AND PRECAUTIONS (5) section, Risk of Aortic Aneurysm and Dissection (5.8) subsection**
- **USE IN SPECIFIC POPULATIONS (8) section, Geriatric Use (8.5) subsection.**
- **PATIENT COUNSELING INFORMATION (17)**
- **MEDICATION GUIDE**

In addition, minor editorial revisions have been made throughout the labeling to reflect these changes.

## **APPROVAL & LABELING**

We have completed our review of these supplemental applications, as amended during the discussion period. They are 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 (Medication Guide), 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 that include labeling changes 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 Microsoft Word format, that includes the changes approved in this supplemental application, as well as annual reportable changes. To facilitate review of your submission(s), 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, call Fariba Izadi, PharmD, Acting Safety Project Manager, at (301) 796-0563.

Sincerely,

*{See appended electronic signature page}*

Joseph G. Toerner, M.D., M.P.H.  
Deputy Director for Safety  
Division of Anti-Infective Products  
Office of Antimicrobial Products  
Center for Drug Evaluation and Research

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/s/  
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JOSEPH G TOERNER  
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