



Our STN: BL 125562/0

BLA FILING NOTIFICATION

Cangene Corporation
Attention: Ms. Allison Kennedy
155 Innovation Drive
Winnipeg, MB R3T 5Y3
Canada

Dear Ms. Kennedy:

This letter is in regard to your biologics license application (BLA) submitted under section 351 of the Public Health Service Act.

We have completed an initial review of your application dated July 25, 2014 for Anthrax Immune Globulin (Intravenous) Human to determine its acceptability for filing. Under 21 CFR 601.2(a), we have filed your application today. The review classification for this application is Priority. Therefore, the review goal date is March 25, 2015. This acknowledgment of filing does not mean that we have issued a license nor does it represent any evaluation of the adequacy of the data submitted.

We are reviewing your application according to the processes described in the Guidance for Review Staff and Industry: Good Review Management Principles and Practices for PDUFA Products. Therefore, we have established internal review timelines as described in the guidance, which include the timeframes for FDA internal milestone meetings. We plan to hold our internal mid-cycle review meeting on November 6, 2014. Please be aware that the timelines described in the guidance are flexible and subject to change based on workload and other potential review issues (e.g., submission of amendments). We will inform you of any necessary information requests or status updates following the milestone meetings or at other times, as needed, during the process.

We will contact you regarding your proposed labeling no later than February 23, 2015. If post marketing study commitments (506B) are required, we will contact you no later than February 23, 2015.

We are still assessing if the application will be presented at an Advisory Committee meeting, but will inform you when we have a decision.

At this time, we have not identified any potential review issues. Our filing review is only a preliminary review, and deficiencies may be identified during substantive review of your application. Following a review of the application, we shall advise you in writing of any action we have taken and request additional information if needed.

If you have any questions, please contact the Regulatory Project Manager, LT Thomas J. Maruna, USPHS, MSc, MLS(ASCP) at (240) 402-8454 or thomas.maruna@fda.hhs.gov.

Sincerely yours,

Iliana Valencia, MS
Chief
Regulatory Project Management Staff
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