SUPPLEMENT APPROVAL
June 22, 2018

Our STN: BL 125314/50

Alba Bioscience Limited
Attention: Mr. Robert Dorris
21 Ellen's Glen Road, Edinburgh Midlothian
Edinburgh EH 17 7QT
United Kingdom

Dear Mr. Dorris:

We have approved your request dated August 18, 2017, to supplement your Biologics License Application (BLA) submitted under section 351(a) of the Public Health Service Act (42 U.S.C. 262) for Blood Grouping Reagent, Anti-D (Monoclonal Blend), to include the manufacture of Albaclone® Anti-D *fusion* at your facility located in Edinburgh, Scotland, United Kingdom. Albaclone® Anti-D *fusion* is for the in vitro detection and identification of human RhD blood group status in patient and donor samples by direct agglutination and indirect antiglobulin test using tube techniques.

We hereby approve the draft package insert labeling submitted under amendment 2, dated April 9, 2018, and the draft carton and container labeling submitted with the supplement, dated August 18, 2017. This is a reminder that as of September 24, 2014, medical devices that are licensed under the PHS Act are subject to certain provisions of the final Unique Device Identifier (UDI) rule. These provisions include the requirement to provide a UDI on the device label and packages (21 CFR 801.20), format dates on the device label in accordance with 21 CFR 801.18, and submit data to the Global Unique Device Identification Database (GUDID) (21 CFR 830 Subpart E). Additionally, please identify each device identifier implemented for the subject device, and the device identifiers that have been discontinued for the subject device as a labeling change in an annual report consistent with 21 CFR 601.12(f)(3). For more information on these requirements, please see the UDI website, http://www.fda.gov/udi.

Please submit all final printed labeling as PDF electronic copy (eCopy) at the time of use and include implementation information on Form FDA 356h as appropriate.

Two draft copies of the proposed introductory advertising or promotional labeling may be voluntarily submitted for advisory comment with a completed Form FDA 2253 to the Advertising and Promotional Labeling Branch at the following address:

Food and Drug Administration
Center for Biologics Evaluation and Research
Document Control Center
10903 New Hampshire Ave.
WO71–G112
Silver Spring, MD 20993-0002

We will include information contained in the above-referenced supplement in your BLA file.

Sincerely,

Orieji Illoh, MD
Director
Division of Blood Components and Devices
Office of Blood Research and Review
Center for Biologics Evaluation and Research