

**SUPPLEMENT APPROVAL**  
**July 8, 2019**

Alba Bioscience Limited  
 Attention: Mr. Robert Dorris  
 5 James Hamilton Way, Allan-Robb Campus  
 Milton Bridge  
 Penicuik, EH 26 OBF  
 United Kingdom

Dear Mr. Dorris:

We have approved your requests submitted March 28, 2019, received March 29, 2019, to supplement your Biologics License Applications (BLA) submitted under section 351(a) of the Public Health Service Act to revise the Instructions for Use of the ORTHO™ Sera Blood Grouping Reagents to include automated testing with ID MTS™ Gel Cards on the ORTHO VISION® Analyzer for the following products:

<b>STN</b>	<b>Name of Biological Products</b>
BL 125304/61*	Blood Grouping Reagent, Anti-D (Monoclonal) (IgM)
BL 125309/56	Blood Grouping Reagent, Anti-N (Murine Monoclonal) (IgG)
BL 125310/56	Blood Grouping Reagent, Anti-Le <sup>a</sup> (Murine Monoclonal)
BL 125311/52	Blood Grouping Reagent, Anti-Le <sup>b</sup> (Murine Monoclonal)
BL 125314/65	Blood Grouping Reagent, Anti-D (Monoclonal Blend)
BL 125342/31	Blood Grouping Reagent, Anti-Fy <sup>b</sup>
BL 125567/8	Blood Grouping Reagent, Anti-Fy <sup>a</sup> (Monoclonal) (IgG)
BL 125568/8	Blood Grouping Reagent, Anti-Jk <sup>a</sup> (Monoclonal)
BL 125569/9	Blood Grouping Reagent, Anti-Jk <sup>b</sup> (Monoclonal)
BL 125570/8	Blood Grouping Reagent, Anti-S (Monoclonal) (IgG)
BL 125571/7	Blood Grouping Reagent, Anti-s (Monoclonal) (IgG)
BI 125572/7	Blood Grouping Reagent, Anti-K (Monoclonal)
BL 125573/7	Blood Grouping Reagent, Anti P1 (Murine Monoclonal)

We hereby approve the draft package insert labeling submitted on July 3, 2019. 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 at <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 supplements in your BLA files.

Sincerely,

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