



Our STN: BL 125300/635

## SUPPLEMENT APPROVAL

GlaxoSmithKline Biologicals  
Attention: Frederick De Brito, Ph.D.  
14200 Shady Grove Road  
VR1500  
Rockville, MD 20850-7464

September 26, 2019

Dear Dr. De Brito:

We have approved your request submitted on March 28, 2019, and received on March 28, 2019, to supplement your Biologics License Application (BLA) under section 351(a) of the Public Health Service Act for Meningococcal (Groups A, C, Y, and W-135) Oligosaccharide Diphtheria CRM197 Conjugate Vaccine (Menveo®) manufactured at your Sovicille (Rosia), Italy facility, to include revisions to the package insert labeling under the *Dosage and Administration* and *How Supplied/Storage and Handling* sections; changes to the design of the outer carton and the container labels for the MenCYW-135 Liquid Conjugate Component and the MenA Lyophilized Conjugate Component; and the addition of linear barcodes to both container labels.

### LABELING

We hereby approve the draft package insert labeling submitted under Amendment 3, dated September 16, 2019, and the draft carton and container labeling submitted under Amendment 2, dated September 4, 2019.

### CONTENT OF LABELING

As soon as possible, but no later than 14 days from the date of this letter, please submit the final content of labeling (21 CFR 601.14) in Structured Product Labeling (SPL) format via the FDA automated drug registration and listing system, (eLIST) as described at <http://www.fda.gov/ForIndustry/DataStandards/StructuredProductLabeling/default.htm>. Information on submitting SPL files using eLIST may be found in the guidance for industry *SPL Standard for Content of Labeling Technical Qs and As* at <http://www.fda.gov/downloads/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/UCM072392.pdf>.

The SPL will be accessible via publicly available labeling repositories.

## **PACKAGE AND CONTAINER LABELS**

Please electronically submit final printed package and container labels that are identical to the package and container labels submitted on September 4, 2019, according to the guidance for industry *Providing Regulatory Submissions in Electronic Format — Certain Human Pharmaceutical Product Applications and Related Submissions Using the eCTD Specifications* at <https://www.fda.gov/regulatory-information/search-fda-guidance-documents/providing-regulatory-submissions-electronic-format-certain-human-pharmaceutical-product-applications>.

All final labeling should be submitted as Product Correspondence to this BLA 125300 at the time of use (prior to marketing) and include implementation information on Form FDA 356h.

## **ADVERTISING AND PROMOTIONAL LABELING**

You may submit two draft copies of the proposed introductory advertising and promotional labeling with 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

You must submit copies of your final advertising and promotional labeling at the time of initial dissemination or publication, accompanied by Form FDA 2253 (21 CFR 601.12(f)(4)).

All promotional claims must be consistent with and not contrary to approved labeling. You should not make a comparative promotional claim or claim of superiority over other products unless you have substantial evidence or substantial clinical experience to support such claims (21 CFR 202.1(e)(6)).

Please submit an amendment to all pending supplemental applications for this BLA that include revised labeling incorporating a revised content of labeling that includes these changes.

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

Sincerely,

Doran L. Fink, M.D., Ph.D.  
Deputy Director - Clinical  
Division of Vaccines and  
Related Products Applications  
Office of Vaccines  
Research and Review  
Center for Biologics  
Evaluation and Research