



## SUPPLEMENT APPROVAL

Our STN: BL 125408/273

Seqirus, Inc.  
Attention: Christina Cocciardo  
50 Hampshire Street  
Cambridge, MA 02139

July 20, 2018

Dear Ms. Cocciardo:

We have approved your request dated April 3, 2018, to supplement your Biologics License Application submitted under section 351(a) of the Public Health Service Act (42 U.S.C. 262) for Influenza Vaccine (Flucelvax® Quadrivalent) manufactured at your Holly Springs, NC facility to include the 2018-2019 United States formulation and associated labeling revisions.

### LABELING

We hereby approve the draft package insert labeling submitted under Amendment 7 dated July 19, 2018, and the draft carton and container labeling submitted April 3, 2018.

Please provide your final content of labeling including the carton and container labels in Structured Product Labeling (SPL) format. All final labeling should be submitted as Product Correspondence to BLA 125408 at the time of use (prior to marketing) and include implementation information on Form FDA 356h.

In addition, please submit the final content of labeling (21 CFR 601.14) in 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>.

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 biologics license application file.

Sincerely yours,

Jerry P. Weir, Ph.D.  
Director  
Division of Viral Products  
Office of Vaccines  
Research and Review  
Center for Biologics  
Evaluation and Research