DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration
Silver Spring MD 20993

NDA 020872/S-038

SUPPLEMENT APPROVAL

sanofi-aventis US LLC
Attention: Doris Sincak
Manager, North America and Global Regulatory Affairs
55 Corporate Drive
Bridgewater, NJ 08807

Dear Ms. Sincak:

Please refer to your Supplemental New Drug Application (sNDA) dated and received November 5, 2015, and your amendments, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for Allegra® Allergy 12 HR (fexofenadine hydrochloride) tablets, 60 mg; Allegra® Allergy 24 HR (fexofenadine hydrochloride) tablets, 180 mg; and Allegra® Allergy 24 HR (fexofenadine hydrochloride) tablets (gelcaps), 180 mg.

This "Prior Approval" supplemental new drug application proposes changes to the graphic design and text of the immediate container (bottle) labels, carton labels, blister carton labels and stretch card (front and back card) labels.

We have completed our review of this application, as amended. It is approved, effective on the date of this letter, for use as recommended in the agreed-upon labeling text.

LABELING

Submit Final Printed Labeling (FPL), as soon as they are available, but no more than 30 days after they are printed. The FPL must be identical to enclosed labeling described below and must be in the "Drug Facts" format (21 CFR 201.66), where applicable.

Allegra® Allergy 12 HR tablets, 60 mg:

- 12-count blister submitted 4/21/16
- 12-count carton submitted 4/21/16, representative of the 24-count carton

Allegra® Allergy 24 HR tablets, 180 mg

- 5-count blister submitted 4/21/16
- 5-count carton submitted 4/21/16, representative of the 15-count carton
- 30-count bottle submitted 11/5/15, representative of the 45- and 70-count bottle
- 30-count carton submitted 4/21/16, representative of the 45- and 70-count carton
- 90-count front and back card submitted 4/21/16

Allegra® Allergy 24 HR tablets (gelcaps), 180 mg

- 8-count blister, submitted on 4/21/16
- 8-count carton submitted on 4/21/16
- 24-count bottle, submitted on 11/5/15, representative of the 40- and 60-count bottle
- 24-count carton submitted 4/21/16, representative of the 40- and 60-count carton
- 80-count front and back card submitted 4/21/16

Submit FPL for all labeling and package configurations approved with NDA 020872/S-038. The FPL should be submitted electronically according to the guidance for industry titled "Providing Regulatory Submissions in Electronic Format – Human Pharmaceutical Product Applications and Related Submissions Using the eCTD Specifications (June 2008)." Alternatively, you may submit 12 paper copies, with 6 of the copies individually mounted on heavy-weight paper or similar material. For administrative purposes, designate this submission "**Final Printed Labeling for approved NDA020872/S-038**." Approval of this submission by FDA is not required before the labeling is used.

DRUG REGISTRATION AND LISTING

All drug establishment registration and drug listing information is to be submitted to FDA electronically, via the FDA automated system for processing structured product labeling (SPL) files (eLIST). At the time that you submit your final printed labeling (FPL), the content of labeling (Drug Facts) should be submitted in SPL format 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 titled "SPL Standard for Content of Labeling Technical Qs and As" at http://www.fda.gov/downloads/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/UCM072392.pdf. In addition, representative container or carton labeling, whichever includes Drug Facts, (where differences exist only in the quantity of contents statement) should be submitted as a JPG file.

REQUIRED PEDIATRIC ASSESSMENTS

Under the Pediatric Research Equity Act (PREA) (21 U.S.C. 355c), all applications for new active ingredients, new indications, new dosage forms, new dosing regimens, or new routes of administration are required to contain an assessment of the safety and effectiveness of the product for the claimed indication(s) in pediatric patients unless this requirement is waived, deferred, or inapplicable.

Because none of these criteria apply to your application, you are exempt from this requirement.

REPORTING REQUIREMENTS

We remind you that you must comply with reporting requirements for an approved NDA (21 CFR 314.80 and 314.81).

If you have any questions, call Sherry Stewart, Regulatory Project Manager, at (301) 796-9618.

Sincerely,

{See appended electronic signature page}

Theresa Michele, MD
Director
Division of Nonprescription Drug Products
Office of Drug Evaluation IV
Center for Drug Evaluation and Research

ENCLOSURES:
Carton and Container Labeling

This is a representation of an electronic record that was signed electronically and this page is the manifestation of the electronic signature.

/s/

THERESA M MICHELE
05/10/2016