



NDA 21920/S-023

SUPPLEMENT APPROVAL

Bionpharma Inc.
Attention: Usha Sankaran
Associate Vice President, Regulatory Affairs
600 Alexander Road
Suite 2-4B
Princeton, NJ 08540

Dear Ms. Sankaran:

Please refer to your Supplemental New Drug Application (sNDA) dated and received August 27, 2018, and your amendments, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (FDCA) for naproxen sodium capsule, 220 mg.

This “Prior Approval” supplemental new drug application provides labeling for two versions of 160-count cartons, to accommodate different distributors. Each version will contain two 80-count bottles.

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, as soon as they are available, but no more than 30 days after they are printed. The final printed labeling must be identical to the labeling below and must be in the “Drug Facts” format (21 CFR 201.66), where applicable.

Submitted Labeling for Approval	Date Submitted
Naproxen Sodium Capsules 160-ct carton (twin-pack (2 x 80-ct)) with “compare to the active ingredient of ALEVE® Liquid Gels” statement	January 22, 2019
Naproxen Sodium Capsules 160-ct carton (twin-pack (2 x 80-ct)) without “compare to the active ingredient of ALEVE® Liquid Gels” statement	January 22, 2019

The final printed labeling should be submitted electronically according to the guidance for industry titled *Providing Regulatory Submissions in Electronic Format — Certain Human Pharmaceutical Product Applications and Related Submissions Using the eCTD Specifications (April 2017, Revision 4)*. For administrative purposes, designate this submission “**Final Printed Labeling for approved NDA 21920/S-023.**” 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 are 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.

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 Tinya Sensie, Regulatory Project Manager at (240) 402-4230.

Sincerely,

{See appended electronic signature page}

Karen Murry Mahoney, MD, FACE
Deputy Director
Division of Nonprescription Drug Products
Office of Drug Evaluation IV
Center for Drug Evaluation and Research

ENCLOSURE(S):
Carton and Container Labeling

This is a representation of an electronic record that was signed electronically. Following this are manifestations of any and all electronic signatures for this electronic record.

/s/

KAREN M MAHONEY
02/20/2019 05:05:43 PM