DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
Rockville, MD 20857

NDA 22-155/S-006

McNeil Consumer Healthcare
Attention: Terry Chan
Senior Associate, Global Regulatory Affairs
7050 Camp Hill Road
Fort Washington, PA 19034-2299

Dear Mr. Chan:

Please refer to your supplemental new drug application dated November 6, 2008, received November 7, 2008, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Children's Zyrtec® Allergy, (1 mg/ mL cetirizine HCl) syrup.

We acknowledge receipt of your submission dated November 12, 2008 and your amendment dated February 23, 2009 that provided revised draft labeling.

This supplemental new drug application provides for:

1. An additional container closure system for the grape flavored, sugar-free formulation of Children's Zyrtec® Allergy syrup to deliver a single 5mg dose.
2. The addition of content uniformity testing for the release of product packaged in the unit dose spoon.

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

The final printed labeling (FPL) must be identical to the revised draft carton and pouch labels for the Children's Zyrtec® Allergy syrup, grape sugar-free dye-free flavor, submitted on February 23, 2009. The FPL must be submitted when available and must be formatted in accordance with the requirements of 21 CFR 201.66 where applicable.

Please submit an electronic version of the FPL according to the guidance for industry titled *Providing Regulatory Submissions in Electronic Format - NDA.* Alternatively, you may submit 20 paper copies of the FPL as soon as it is available but no more than 30 days after it is printed. Individually mount 15 of the copies on heavy-weight paper or similar material. For administrative purposes, designate this submission "**FPL for approved supplement NDA 22-155/S-006**." Approval of this submission by FDA is not required before the labeling is used.

We remind you that the flags "New Flavor" and "NEW! pre-filled dosing spoon" on the principal display panel of each SKU should be removed from the principal display panel after six months of marketing.

If you issue a letter communicating important information about this drug product (i.e., a "Dear Health Care Professional" letter), we request that you submit a copy of the letter to this NDA and a copy to the following address:

MEDWATCH
Food and Drug Administration
Suite 12B05
5600 Fishers Lane
Rockville, MD 20857

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 Janice Adams-King, Regulatory Project Manager, at (301) 796-3713.

Sincerely,

{See appended electronic signature page}

Joel Schiffenbauer, M.D.
Deputy Director
Division of Nonprescription Clinical Evaluation
Office of Nonprescription Products
Center for Drug Evaluation and Research

Enclosure