



NDA 19-717/S-052

Eli Lilly and Company
Attention: Jeffrey L. Winn, D.D.S., R.Ph.
Senior Regulatory Research Scientist
US Regulatory Affairs
Lilly Corporate Center
Indianapolis, IN 46285

Dear Dr. Winn:

Please refer to your supplemental new drug application dated February 10, 2003, received February 11, 2003, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Humulin® 70/30 (70% human insulin isophane suspension and 30% human insulin injection, [rDNA origin]).

We acknowledge receipt of your submission dated April 9, 2003.

This “Changes Being Effected” supplemental new drug application provides for an addition of tamper evident tape with the text “**If the seal is broken before first use, contact pharmacist**” added to the 3 mL disposable insulin delivery devices (HP 8770, Humulin 70/30 Pen) carton label.

We completed our review of this supplemental new drug application. It is approved, effective on the date of this letter, for use as recommended in the final printed labeling (FPL) submitted on April 9, 2003.

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, HFD-410
FDA
5600 Fishers Lane
Rockville, MD 20857

NDA 19-717/S-052

Page 2

We remind you that you must comply with the requirements for an approved NDA set forth under 21 CFR 314.80 and 314.81.

If you have any questions, call Julie Rhee, Regulatory Project Manager, at (301) 827-6424.

Sincerely,

{ See appended electronic signature page }

David G. Orloff, M.D.
Director
Division of Metabolic
and Endocrine Drug Products, HFD-510
Office of Drug Evaluation II
Center for Drug Evaluation and Research

Enclosure: Carton label (circular SH 8732 FSAMS) submitted on April 9, 2003

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

/s/

David Orloff

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