

Food and Drug Administration Rockville, MD 20857

NDA 20-868/S-006

Pfizer, Inc.

Attention: Pritpal Nijjar

Regulatory Manager

235 East 42nd St. New York, NY 10017

Dear Ms. Nijjar:

Please refer to your supplemental new drug application dated October 9, 2003, received October 17, 2003, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for FLAGYL[®] ER (metronidazole).

We acknowledge receipt of your submission dated October 30, 2003.

This "Changes Being Effected" supplemental new drug application provides for revised labeling to comply with the Final Rule entitled "**Labeling Requirements for Systemic Antimicrobial Drug Products Intended for Human Use"** (68FR 6062, February 6, 2003) and respond to our CBE request letter dated September 11, 2003.

This "Changes Being Effected" supplemental new drug application provides for the following additions to the package insert:

Location	Text
At the beginning	To reduce the development of drug-resistant bacteria and maintain the
of the label, under	effectiveness of FLAGYL ER and other antibacterial drugs, FLAGYL ER
"PRODUCT	should be used only to treat or prevent infections that are proven or strongly
NAME"	suspected to be caused by bacteria.
INDICATIONS	To reduce the development of drug-resistant bacteria and maintain the
AND USAGE	effectiveness of FLAGYL ER and other antibacterial drugs, FLAGYL ER
	should be used only to treat or prevent infections that are proven or strongly
	suspected to be caused by susceptible bacteria. When culture and susceptibility
	information are available, they should be considered in selecting or modifying
	antibacterial therapy. In the absence of such data, local epidemiology and
	susceptibility patterns may contribute to the empiric selection of therapy.
PRECAUTIONS	Prescribing FLAGYL ER in the absence of a proven or strongly suspected
section, under	bacterial infection or a prophylactic indication is unlikely to provide benefit to
"General"	the patient and increases the risk of the development of drug-resistant
subsection	bacteria."
PRECAUTIONS	Patients should be counseled that antibacterial drugs including FLAGYL ER
section, under	should only be used to treat bacterial infections. They do not treat viral
"Information for	infections (e.g., the common cold). When FLAGYL ER is prescribed to treat a
Patients"	bacterial infection, patient should be told that although it is common to feel

better early in the course of therapy, the medication should be taken exactly as
directed. Skipping doses or not competing the full course of therapy may (1)
decrease the effectiveness of the immediate treatment and (2) increase the
likelihood that bacterial will develop resistance and will not be treatable by
FLAGYL ER or other antibacterial drugs in the future.

We have completed the review of this supplemental application, as amended, and have concluded that adequate information has been presented to demonstrate that the drug product is safe and effective for use as recommended in the final printed label submitted on October 30, 2003 (enclosed). Accordingly, this supplemental application is approved effective on the date of this letter.

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

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 Robin Anderson, R.N., M.B.A., Labeling Reviewer at (301) 827-2127.

Sincerely,

{See appended electronic signature page}

Renata Albrecht, M.D.
Director
Division of Special Pathogen and Immunologic
Drug Products
Office of Drug Evaluation IV
Center for Drug Evaluation and Research

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/s/

Renata Albrecht 3/17/04 10:58:43 PM