



NDA 12-623/S-059

Pfizer, Inc.  
Attention: Pritpal Nijjar  
Regulatory Manager  
235 East 42<sup>nd</sup> 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<sup>®</sup> (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:

<b>Location</b>	<b>Text</b>
At the beginning of the label, under “ <b>PRODUCT NAME</b> ”	To reduce the development of drug-resistant bacteria and maintain the effectiveness of FLAGYL and other antibacterial drugs, FLAGYL should be used only to treat or prevent infections that are proven or strongly suspected to be caused by bacteria.
<b>INDICATIONS AND USAGE</b>	To reduce the development of drug-resistant bacteria and maintain the effectiveness of FLAGYL and other antibacterial drugs, FLAGYL 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.
<b>PRECAUTIONS</b> section, under “General” subsection	Prescribing FLAGYL in the absence of a proven or strongly suspected bacterial infection or a prophylactic indication is unlikely to provide benefit to the patient and increases the risk of the development of drug-resistant bacteria.”
<b>PRECAUTIONS</b> section, under “Information for	Patients should be counseled that antibacterial drugs including FLAGYL should only be used to treat bacterial infections. They do not treat viral infections (e.g., the common cold). When FLAGYL 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 completing 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 or other antibacterial drugs in the future.
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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/

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Renata Albrecht  
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