



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
Rockville, MD 20857

May 24, 2004

FILE COPY

Mr. Robert W. Pollock
Lachman Consultant Services, Inc.
1600 Stewart Avenue
Suite 604
Westbury, New York 11590

Dear Mr. Pollock:

Your petition requesting the Food and Drug Administration to make a determination that Oxycodone and Acetaminophen Tablet, USP combination drug products, in strengths of 2.5 mg/400 mg, 5 mg/ 400 mg, 7.5 mg/400 mg and 10 mg/ 400 are suitable for submission in an ANDA, was received by this office on 05/24/2004. It was assigned docket number 2004P-0243 /CP 1 and it was filed on 05/24/2004. Please refer to this docket number in future correspondence on this subject with the Agency.

Please note that the acceptance of the petition for filing is a procedural matter in that in no way reflects an agency decision on the substantive merits of the petition.

Sincerely,

Lyle D. Jaffe
Division of Dockets Management
Office of Management Programs
Office of Management

2004P-0243

ACK1