



Food and Drug Administration  
Rockville, MD 20857

May 14, 2004

Kathleen M. Sanzo  
Morgan Lewis & Bocklus LLP  
1111 Pennsylvania Avenue, NW  
Washington, DC 20004

Dear Ms. Sanzo:

Your petition, submitted on behalf of Pfizer Inc, requesting the Food and Drug Administration to deny approval of a New Drug Application 21-426 for Omnitrop 5.8 mg somatropin [ rDNA origin ] for injection, lyophilized powder and diluent with preservative was received by this office on 05/14/2004. It was assigned docket number 2004P-0231/CP1 and it was filed on 05/14/2004. 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 it in no way reflects an agency decision on the substantive merits of the petition.

Sincerely,

Jennie C. Butler, Director  
Division of Dockets Management  
Office of Management Programs  
Office of Management

2004P.0231

ACK 1