DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
Rockville MD 20857

August 7, 2006

FILE COPY

David L. Rosen, B.S. Pharm, J.D.
Foley & Lardner LLP
3000 K Street, N.W., Suite 500
Washington, DC 20007-5143

Dear Mr. Rosen:

Your petition requesting the Food and Drug Administration to investigate and take regulatory action as necessary and appropriate to protect surgical patients from a potential significant safety risk in connection with Propofol Injectable Emulsion marketed by Bedford Laboratories was received by this office on 08/07/2006. It was assigned docket number 2006P-0311/CP 1 and it was filed on 08/07/2006. 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,

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

2006P-0311

ACK 1