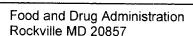


Public Health Service



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

20067-0311

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