DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
Rockville, MD 20857

July 3, 2006

FILE COPY

Mr. Micheal C. Beckloff
Beckloff Associates, Inc.
7400 West 110th Street
Suite 300
Overland Park, Kansas 66210

Dear Mr. Beckloff:

Your petition requesting the Food and Drug Administration to declare that the drug products containing 0.9% Sodium Chloride Injection, USP, in 3-, 5-, 15-, and 30-mL volumes (strengths), is suitable for submission as an abbreviated new drug application, was received by this office on 07/03/2006. It was assigned docket number 2006P-0272/CP1 and it was filed on 07/03/2006. 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,

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

2006P-0272

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