## DEPARTMENT OF HEALTH & HUMAN SERVICES

Food and Drug Administration Silver Spring MD 20993

September 18, 2019

Kurt R. Karst Hyman, Phelps & McNamara, P.C. 700 Thirteenth Street, N.W., Suite 1200 Washington, DC 20005-5929

Sent via email to: kkarst@hpm.com

Dear Petitioners:

Your petition to the Commissioner of Food and Drug Administration requesting to determine that Cefazolin for Injection USP, 2 g vial – administered via intramuscular injection, intravenous direct (bolus) injection, and intravenous infusion is suitable for submission in an Abbreviated New Drug Application was received by this office on 09/17/2019.

It was assigned docket number FDA-2019-P-4363. 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,

Dynna Bigby Supervisory Administrative Proceedings Specialist Dockets Management Staff FDA/Office of Operations (OO)