DEPARTMENT OF HEALTH & HUMAN SERVICES

Food and Drug Administration
Rockville, MD 20857

January 27, 2020

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

Sent via email to: kkarst@hpm.com

Dear Petitioner:

Your petition to the Commissioner of Food and Drug Administration requesting to determine whether Micro-K LS Packets (potassium chloride) 20mEq Extended-release Liquid Suspension, approved under New Drug Application (NDA) number 019561 has been voluntarily withdrawn for reasons of safety or effectiveness was received by this office on 01/27/2020.

It was assigned docket number FDA-2020-P-0438. 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

Digitally signed by
Dynna Bigby
Date: 2020.01.27
14:51:37 -05'00'

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