



September 13, 2022

Kurt R. Karst
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Washington, DC 20005-5929

Sent via email to: kkarst@hpm.com

Dear Petitioner:

Your submission requesting that the Commissioner of Food and Drug determine whether Hydrochlorothiazide Oral Solution, 50 mg/5mL, approved under Abbreviated New Drug Application number 088587, held by Roxane Laboratories, Inc., has been voluntarily withdrawn for reasons of safety or effectiveness was received and processed under CFR 10.30 by this office on 09/13/2022.

It was assigned docket number FDA-2022-P-2229. Please refer to this docket number in future correspondence on this subject with the Agency.

Please note, the acceptance of the petition for filing is a procedural matter and in no way reflects the Agency's decision on the substantive merits of the petition.

Sincerely,

Karen Malvin
Acting Director
Dockets Management Staff
FDA/Office of Operations (OO)

cc: inamata@hpm.com