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VIA REGULATIONS.GOV

Division of Dockets Management (HFA-305) Food and Drug Administration Department of Health and Human Services 5360 Fishers Lane, Room 1061 Rockville, MD 20852

RE: Withdrawal of Petition; Docket No. FDA-2020-P-1849

Dear Sir or Madam:

This Petitioner hereby requests withdrawal of the above-referenced citizen petition requesting that FDA designate a new reference standard for hydrocortisone and neomycin sulfate and polymyxin B sulfate otic suspension/drops, 1%; Eq 3.5mg Base/mL; 10,000 Units/mL.

Sincerely yours,

DAVILL Rosen

David L. Rosen, B.S. Pharm., JD

Cc: Alvin Chi, J.D.

Regulatory Counsel

Center for Drug Evaluation and Research

Office of Regulatory Policy Sungjoon.Chi@fda.hhs.gov