December 7, 2020

**VIA ELECTRONIC SUBMISSION**

Division of Dockets Management
Food and Drug Administration (HFA-305)
Department of Health and Human Services
5630 Fishers Lane, Room 1061
Rockville, MD 20852

**CITIZENS PETITION – WITHDRAWAL REQUEST**

**RE: FDA-2019-P-4879**

Dear Sir or Madam:

Lachman Consultant Services, Inc. hereby requests to withdraw the submitted Citizen Petition dated October 18, 2019; assigned Docket number FDA-2019-P-4879.

The Citizen Petition was submitted pursuant to the Federal Food, Drug and Cosmetic Act ("FDC Act") and in accordance with 21 C.F.R. §§ 10.25(a) and 10.30 to request that FDA designate Nitrofurantoin Oral Suspension, 25 mg/5 mL, approved under abbreviated new drug application (ANDA) 201355 held by Nostrum Laboratories Inc., as a reference standard in the *Approved Drug Products with Therapeutic Equivalence Evaluations* (the Orange Book) (Docket No. FDA-2019-P-4879).

Please do not hesitate to contact the undersigned if additional information is required in order to complete withdrawal of FDA-2019-P-4879 as the Lachman consultant who originally submitted the petition is no longer with the firm.

Sincerely,

D.Sloane@LachmanConsultants.com
Digitally signed by D.Sloane@LachmanConsultants.com
Date: 2020.12.07 10:49:22 -05'00'

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