

March 18, 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-2020-P-0598**

Dear Sir or Madam:

Lachman Consultant Services, Inc. hereby requests to withdraw the Citizen Petition submitted on January 22, 2020; assigned Docket number FDA-2020-P-0598.

The Citizen Petition was submitted pursuant to the Federal Food, Drug and Cosmetic Act (“FD&C Act”) and in accordance with 21 CFR § 10.25(a) and 10.30 requesting the Commissioner of the Food and Drug Administration to designate a suitable alternative reference standard (RS) for the purpose of conducting *in vivo* bioequivalence studies to support an ANDA application for Prochlorperazine Maleate Tablets USP, 10 mg.

Please advise if additional information is required in order to complete withdrawal request.

Sincerely,

Michelle R. Ryder  
Principal Consultant  
Lachman Consulting Services, Inc.