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VIA REGULATIONS.GOV

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

RE: Withdrawal of Suitability Petition; Docket No. FDA-2020-P-1252

Dear Sir or Madam:

This Petitioner hereby requests withdrawal of the above-referenced citizen petition requesting that the Commissioner of Food and Drugs allow the submission of an ANDA for Lidocaine Hydrochloride Injection USP, 20mg /2mL (10mg/mL), 2 mL Fill vials (total vial content 20mg) pursuant to Section 505(j)(2)(C) of the Federal Food, Drug, and Cosmetic Act and 21 CFR § 314.93.

Sincerely yours,

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