Food and Drug Administration Silver Spring MD 20993

February 3, 2022

Michelle Ryder Lachman Consultant Services, Inc. 1600 Stewart Avenue, Suite 604 Westbury, NY 11590

Sent via email to: m.ryder@lachmanconsultants.com

Dear Petitioner:

Your submission requesting that the Commissioner of Food and Drug Administration declare that Diazepam Injection USP, 20 mg/4 mL (5 mg/mL) single dose prefilled syringe is suitable for submission as an ANDA was received and processed under CFR 10.30 by this office on 02/02/2022.

It was assigned docket number FDA-2022-P-0118. Please refer to this docket number in future correspondence on this subject with the Agency.

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

Sincerely,

Dynna Bigby Supervisory Administrative Proceedings Officer Dockets Management Staff FDA/Office of Operations (OO)