

DEPARTMENT OF HEALTH AND HUMAN SERVICES FOOD AND DRUG ADMINISTRATION		
<small>DISTRICT ADDRESS AND PHONE NUMBER</small> 1431 Harbor Bay Parkway Alameda, CA 94502-7070 (510) 337-6700 Fax: (510) 337-6702	<small>DATE(S) OF INSPECTION</small> 12/10/2018-12/19/2018*	
<small>NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED</small> Christopher F. Zuccarelli, Chief Operating Officer		
<small>FIRM NAME</small> Leiter's Enterprises, Inc.	<small>STREET ADDRESS</small> 17 Great Oaks Blvd	
<small>CITY, STATE, ZIP CODE, COUNTRY</small> San Jose, CA 95119-1359	<small>TYPE ESTABLISHMENT INSPECTED</small> Outsourcing Facility	
This document lists observations made by the FDA representative(s) during the inspection of your facility. They are inspectional observations, and do not represent a final Agency determination regarding your compliance. If you have an objection regarding an observation, or have implemented, or plan to implement, corrective action in response to an observation, you may discuss the objection or action with the FDA representative(s) during the inspection or submit this information to FDA at the address above. If you have any questions, please contact FDA at the phone number and address above.		
<b>DURING AN INSPECTION OF YOUR FIRM I OBSERVED:</b> <b>OBSERVATION 1</b>  The responsibilities and procedures applicable to the quality control unit are not in writing and fully followed.  Specifically,  Your firm's management of the building monitoring system alarms lacks adequate quality control review. There are no time frames established in your firm's procedures to ensure alarms are reviewed in a timely manner and quality control is not required to review alarm acknowledgements to ensure alarms are handled appropriately and do not impact drug products.		
<b>OBSERVATION 2</b> Employees engaged in the processing of a drug product lack the training required to perform their assigned functions.  Specifically,  During routine visual inspections of filled drug products, an operator is not required to take a break unless they have been inspecting for (b) (4). An operator may inspect (b) (4) units for (b) (4) before taking a (b) (4) break. Your firm		
<b>SEE REVERSE OF THIS PAGE</b>	<small>EMPLOYEE(S) SIGNATURE</small> Ashar P Parikh, Investigator	<small>DATE ISSUED</small> 12/19/2018  <small>Ashar P Parikh                      Investigator                      Signed by Ashar P. Parikh -S                      Date Signed 12-19-2018 13:09:08</small> X

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<p>currently qualifies the visual inspectors if they can pass (b) (4) visual inspection qualifications of (b) (4) identifying at least (b) (4) % of the units with defects. This is inadequate to ensure an operator is capable of detecting drug product units with defects visible particulates during the entire (b) (4) visual inspection process.</p>		
<p><b>OBSERVATION 3</b>                      Aseptic processing areas are deficient regarding the system for cleaning and disinfecting the equipment to produce aseptic conditions.</p> <p>Specifically,</p> <p>Your firm does not routinely clean the wheels of chairs and rolling carts in your ISO 7 cleanroom. The carts are moved throughout the cleanroom during operations as observed during aseptic processing on December 12, 2018. Your firm's SOP 3.020 "Cleaning and Disinfection of ISO 7 and ISO 8 Areas" has no requirement for the technicians to clean the chair and cart wheels.</p>		
<p><b>*DATES OF INSPECTION</b>                      12/10/2018(Mon), 12/11/2018(Tue), 12/12/2018(Wed), 12/13/2018(Thu), 12/14/2018(Fri),                      12/19/2018(Wed)</p>		
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