May 5, 2020

UPS NEXT DAY
SIGNATURE REQUIRED

Lawrence J. Frieders  
Owner and Pharmacist in Charge  
Techni Med Inc. dba The Compounder 
340 Marshall Avenue, Unit 100  
Aurora, IL 60506-5649

Dear Mr. Frieders:

We are enclosing a copy of the Establishment Inspection Report (EIR) for the inspection conducted at your facility, Techni Med Inc. dba The Compounder, located at 340 Marshall Avenue, Unit 100, Aurora, IL 60506-5649, from September 23 to October 10, 2019, by the U.S. Food and Drug Administration (FDA). In addition, we are enclosing the letter sent to the Illinois Department of Financial and Professional Regulation, Division of Professional Regulation, Illinois State Board of Pharmacy for follow up.

When the Agency considers an inspection to be “closed” under 21 C.F.R. 20.64(d)(3), it will release a copy of the EIR to the inspected establishment.

The Agency continually works to make its regulatory process and activities more transparent for regulated industry. Releasing this EIR to you is part of this effort. The copy being provided to you comprises the narrative portion of the report; it may reflect redactions made by the Agency in accordance with the Freedom of Information Act (FOIA) and 21 C.F.R. Part 20. This, however, does not preclude you from requesting and possibly obtaining any additional information under FOIA.

If there is any question about the released information, please contact Brian Garthwaite, Compliance Officer at (612) 758-7132 or by email at: ORAPHARM3_RESPONSES@fda.hhs.gov.

Sincerely,

Nicholas F. Lyons  
Director Compliance Branch 
Division of Pharmaceutical Quality Operations III

Enclosure: Redacted 1st party EIR and State Referral Letter