FDA U.S. FOOD & DRUG ADMINISTRATION
OFFICE OF REGULATORY AFFAIRS
OFFICE OF PHARMACEUTICAL QUALITY OPERATIONS

May 20, 2020

Case #: 604605

VIA ELECTRONIC MAIL

Christopher S. Gilbert, PharmD, Owner
People's Custom Rx and Clinical Care, LLC
785 E. Brookhaven Circle East
Memphis, TN 38117-4501
cgilbert@peoplescustomrx.com

Dear Dr. Gilbert:

We are enclosing a copy of the Establishment Inspection Report (EIR) for the inspection conducted at your facility, People's Custom Rx and Clinical Care, LLC, 785 Brookhaven Circle East, Memphis, Tennessee 38117, from July 16, 2019, to July 24, 2019, by the U.S. Food and Drug Administration (FDA).

When the Agency concludes that an inspection is "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 you have questions regarding the contents of this letter, you may contact Compliance Officer Rebecca Asente via phone at (504) 846-6104 or email Rebecca.asente@fda.hhs.gov.

Sincerely,

John W. Diehl -S3

Digitally signed by John W. Diehl -S3
DN c US, o U.S. Government, ou HHS, ou FDA, ou People, cn John W. Diehl -S3, 0.9.2342.19200300.100.1.1 2000099727
Date 2020.05.20 14 02 50 -05'00'

John W. Diehl, M.S
Director, Compliance Branch
Office of Pharmaceutical Quality Operations, Division II

U.S. Food & Drug Administration
Office of Pharmaceutical Quality Operations, Division II
4040 N. Central Expressway, Suite 300
Dallas, Texas 75204
www.fda.gov