WARNING LETTER

Date : November 30, 2020

TO: sales@avazo.com

Avazo-Healthcare, LLC
17 Regina Place
Warminster Bucks, PA 18974

605 Louis Dr Suite 502
Warminster, PA 18974

web@cbdmarketweb.com

CBD MarketWeb
605 Louis Dr Suite 02
Warminster, PA 18974

info@covidtestshop

COVID Test Shop
605 Louis Dr Suite 502
Warminster, PA 18974

RE: Adulterated, Unapproved, and Misbranded Products Related to Coronavirus Disease 2019 (COVID-19)

This is to advise you that the United States Food and Drug Administration (FDA) reviewed your websites at the Internet addresses https://www.covidtestshop.com and www.cbdmarketweb.com on August 21, 2020 through November 16, 2020. We have also reviewed your social media websites at https://www.facebook.com/avazodeets/, https://www.facebook.com/cbdmarketweb/, and https://www.instagram.com/cbdmarketweb/, which direct consumers to your websites https://www.covidtestshop.com and www.cbdmarketweb.com to purchase your products. The FDA has observed that your websites offer COVID-19[1] test kit products and CBD products for sale in the United States and that these products are intended to mitigate, prevent, treat, diagnose, or cure COVID-19 in people.

There is currently a global outbreak of respiratory disease caused by a novel coronavirus that has been named "severe acute respiratory syndrome coronavirus 2" (SARS-CoV-2). The disease caused by the virus has been named "Coronavirus Disease 2019" (COVID-19). On January 31, 2020, the Department of Health and Human Services (HHS) issued a declaration of a public health emergency related to COVID-19 and mobilized the Operating Divisions of HHS.[2] In addition, on March 13, 2020, the President

---

[1] As explained in the next paragraph, there is currently an outbreak of a respiratory disease named "Coronavirus Disease 2019" (COVID-19).

[2] Secretary of Health and Human Services Alex M. Azar II, Determination that a Public Health Emergency Exists. Jan. 31, 2020. (Accessible at https://www.phe.gov/emergency/news/healthactions/phe/Pages/2019-nCoV.aspx). The declaration has been renewed for an additional 90 days three times. The most recent renewal went into effect on October 23, 2020. Secretary of Health and Human Services Alex M. Azar II, Renewal of Determination that a Public Health Emergency Exists. October 23, 2020. (Accessible at https://www.phe.gov/emergency/news/healthactions/phe/Pages/covid19-23Oct2020.aspx).

Test Cassette and the Antigen Rapid Test Device are labeled by the "authorized labeling" in accordance with the requirements of their respective EUAs.

For more information about FDA's regulation of devices used to mitigate, prevent, treat, diagnose, or cure COVID-19; frequently asked questions and other helpful resources, website at

Timothy T. Stenzel -