WARNING LETTER

Date: March 6, 2020

TO: partner.services@jimbakkershow.com, website@jimbakkershow.com – James Bakker, The Jim Bakker Show
180 Grace Chapel Rd.
Blue Eye, MO 65611

RE: 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) and the Federal Trade

[from your video entitled "A Close Look At What's Not Being Said About the Coronavirus (Day 1)" at [44:48], https://jimbakkershow.com/watch/?guid=3861]

- "Silver Solution has been proven … to kill every pathogen it has ever been tested on … and it can kill any of these known viruses …"

- "So the virus, like the coronavirus that we're talking about … affects the lung tissue so what you can do … put it straight … in a nebulizer which then creates a steam and you breathe it in and it will go directly into your lungs where that virus is and any other infection"

"You should take immediate action to correct the violations cited in this letter. The violations cited in this letter are not meant to be an all-inclusive list. It is your responsibility to ensure that the products you sell are in compliance with the FD&C Act and FDA's implementing regulations. We advise you to review your websites, product labels, and other labeling and promotional materials to ensure that you are not representing your products for a COVID-19 related use for which they have not been approved by FDA, and that you do not make claims that misbrand the products in violation of the FD&C Act**. Within 48 hours, please send an email to** COVID-19-Task-Force-CFSAN@fda.hhs.gov describing the specific steps you have taken to correct these violations. Include an explanation of each step being taken to prevent the recurrence of violations, as well as copies of related documentation. Failure to immediately correct the violations cited in this letter may result in legal action, including, without limitation, seizure and injunction.

FDA is advising consumers not to purchase or use certain products that have not been approved, cleared, or authorized by FDA and that are being misleadingly represented as safe and/or effective for the treatment or prevention of COVID-19. Your firm will be added to a published list on FDA's website of firms and websites that have received warning letters from FDA concerning the sale or distribution of COVID-19 related products in violation of the FD&C Act. This list can be found at www.fda.gov/consumers/health-fraud-scams/fraudulent-coronavirus-disease-covid-19-products. Once you have taken corrective actions and such actions have been confirmed by the FDA, the published list will be updated to indicate that your firm has taken appropriate corrective action.

If you cannot complete corrective action within 48 hours, state the reason for the delay and the time within which you will complete the corrections. If you believe that your products are not in violation of the FD&C Act, include your reasoning and any supporting information for our consideration.

If you are not located in the United States, please note that products that appear to be misbranded or unapproved new drugs are subject to detention and refusal of admission if they are offered for importation into the United States. We may advise the appropriate regulatory officials in the country from which you operate that FDA considers your product(s) listed above to be unapproved and misbranded products that cannot be legally sold to consumers in the United States. Please direct any inquiries to F e4 (r)3 (y)20 (f)3 (r)3 (om)-(