October 3, 2022

Dustin Bryce, Public Relations
Interest Of Justice
Entrega General, Lista Correos
San Isidro, San Jose, PÉREZ ZELEDÓN
Costa Rica, 11901

Sent via email to: contact@interestofjustice.org

Dear Petitioner:

Your submission requesting that the Commissioner of Food and Drug revoke FDA Authorization for Moderna, Pfizer-BioNTech Bivalent COVID-19 Vaccines for Use as a Booster Dose, and revoke all other authorization for mRNA Gene Therapies misbranded as vaccines which do not confer immunity because FDA January 2020 guidance: "Long Term Follow-Up After Administration of Human Gene Therapy Products Guidance for Industry," states the use of these experimental products affects DNA with an unreasonable risk to use in more than a few people at a time was received and processed under CFR 10.30 by this office on 09/30/2022.

It was assigned docket number FDA-2022-P-2411. Please refer to this docket number in future correspondence on this subject with the Agency.

Please note, the acceptance of the petition for filing is a procedural matter and in no way reflects the Agency's decision on the substantive merits of the petition.

Sincerely,

Karen P. Malvin -S
Digitally signed by Karen P. Malvin -S
Date: 2022.10.03 13:22:54 -04'00'

Karen Malvin
Acting Director
Dockets Management Staff
FDA/Office of Operations (OO)