DEPARTMENT OF HEALTH & HUMAN SERVICES Public Health Service

Food and Drug Administration
Silver Spring MD 20993

June 8, 2020

Paola Brown
President
Americans for Homeopathy Choice Foundation
1601 18th Street NW, Suite 4
Washington, DC 20009

Sent via email to: PaolaBrownEnglish@gmail.com

Dear Petitioner:

Your petition to the Commissioner of Food and Drug Administration requesting the FDA establish regulations that would assure consumers: that drug products labeled homeopathic" are either included in the HPUS or can be reasonably expected to be accepted for inclusion in the HPUS because they meet eligibility thresholds, as determined by relevant third-party review; that products that do not meet the foregoing criteria are not permitted to be labeled "homeopathic;" that homeopathic drugs are free of adulteration and properly labeled; and that FDA applies standards appropriate for low-risk products when evaluating the risks of homeopathic drugs was received by this office on 06/05/2020.

It was assigned docket number FDA-2020-P-1510. Please refer to this docket number in future correspondence on this subject with the Agency.

Please note that the acceptance of the petition for filing is a procedural matter in that it in no way reflects an agency decision on the substantive merits of the petition.

Sincerely,

Dynna Gorham-bigby -S
Digitally signed by Dynna Gorham-bigby -S
Date: 2020.06.08 13:29:10 -04'00'

Dynna Bigby
Supervisory Administrative Proceedings Officer
Dockets Management Staff
FDA/Office of Operations (OO)