January 25, 2022

Via Electronic Submission

Division of Dockets Management
Food and Drug Administration
Department of Health and Human Services
5630 Fishers Lane, Room 1061 (HFA-305) Rockville, MD 20852

Re: Citizen Petition- Requesting FDA-CVM To Publish Publicly On The Public FDA FOIA Reading Room Website, *FDA FOIA Documents Relating To HPP Being A Kill Step For Raw Pet Food Products Under FDA Regulatory Authority,* In Accordance With FOIA Law.

To whom it may concern:

The undersigned submits this petition under the Freedom of Information Act ("FOIA"), 5 U.S.C. § 552, and in accordance with 21 C.F.R. § 10.30(b), requesting both the FDA-CVM and FDA FOIA Department comply with FOIA law and post publicly on the FDA FOIA reading room all documents requested through various FOIA requests pertaining to HPP Being A Kill Step For Raw Pet Food Under FDA Regulatory Authority.

A. Action Requested

Petition requests FDA-CVM to post publicly on the FDA FOIA reading room all documents requested through 3+ FOIA requests pertaining to HPP Being A Kill Step For Raw Pet Food Under FDA Regulatory Authority.

B. Statement of Grounds

FOIA law requires each agency to make available for public inspection, in electronic format, copies of all records that have previously been released under FOIA and "(I) that because of the nature of their subject matter, the agency determines have become or are likely to become the subject of subsequent requests for substantially the same records; or (II) that have been requested 3 or more times." 5 U.S.C. § 552(a)(2)(D).

The following records request:

- *FDA records for or relating to HPP being a kill step for pet food and/or raw pet food*

These records have been requested 3 or more times. FDA-CVM is now required to comply with law U.S.C. § 552(a)(2)(D).

FDA-CVM should provide a final response to this citizen petition within the required 180 days, informing me that FDA-CVM will be complying with this provision of the law, and that FDA-CVM posting these documents on the FDA-CVM FOIA reading room.

C. Environmental Impact

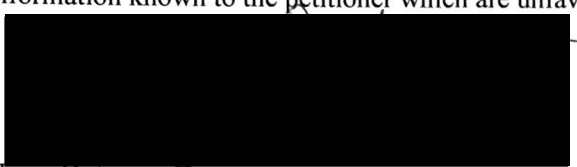
A claim for categorical exclusion of the requirement for submission of an environmental assessment or environmental impact statement is made pursuant to 21 C.F.R. §§ 25.30 and 25.34.

D. Economic Impact

In accordance with 21 C.F.R. § 10.30(b), economic impact information is to be submitted only when requested by FDA following review of the petition. The undersigned will promptly provide this information if requested.

E. Certification

The undersigned certifies, that, to the best knowledge and belief of the undersigned, this petition includes all information and views on which the petition relies, and that it includes representative data and information known to the petitioner which are unfavorable to the petition.



Harry Duty