

Food and Drug Administration Silver Spring MD 20993

February 22, 2022

Harry Duty
(b) (6)

Sent via email to: (b) (6)

Dear Petitioner:

Your submission requesting that the Commissioner of Food and Drug comply with FOIA law and post publicly on the FDA FOIA reading room all documents requested through various FOIA requests pertaining to Testing Records From 2015 Bravo Recall Due to FDA-CVM Regulating By Their Opinion was received and processed under CFR 10.30 by this office on 02/19/2022.

It was assigned docket number FDA-2022-P-0208. Please refer to this docket number in future correspondence on this subject with the Agency.

Also, note that 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,

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