Elizabeth Brehm

From: McNeill, Lorrie <Lorrie.McNeill@fda.hhs.gov>
Sent: Thursday, May 26, 2022 5:07 PM
To: Elizabeth Brehm
Subject: RE: [EXTERNAL] FW: Discrepancies in deaths in Pfizer clinical trial

Dear Ms. Brehm,

We acknowledge your correspondence on behalf of Informed Consent Action Network related to the posted reviews for Comirnaty (COVID-19 vaccine, mRNA), specifically, the Summary Basis of Regulatory Action, Clinical Review memo and Statistical Review memo.

We are unable to respond substantively at this time due to resource constraints and the ongoing pandemic response. It is thus necessary for the Agency to prioritize its responses to individual inquiries. We note that the Agency is providing to your law firm, in response to a FOIA request, an unprecedented volume of records that comprise the license application for this product. Additionally, there are more than 50 pending FOIA requests submitted by your firm for data and information about COVID-19 vaccines.

Sincerely,

Lorrie H. McNeill
Director

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