

Attachment D

OMB Control No. 0910-0497

Exp. Date: 12/31/2026

Paperwork Reduction Act Statement: According to the Paperwork Reduction Act of 1995, an agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a valid OMB control number. The time required to complete this Reminder Letter is estimated to average 1 minute per response, including the time for reviewing instructions and completing and reviewing the collection of information.

Send comments regarding this burden estimate or any other aspects of this collection of information, including suggestions for reducing burden, to PRAStaff@fda.hhs.gov.

This study is being conducted on behalf of the U.S. Food and Drug Administration.

Focus Groups Examining Consumer Reporting of Adverse Events

REMINDER LETTER

Dear [Participant Name]:

This is a reminder letter that the research study about consumers' experiences with prescription and over-the-counter drugs in which you agreed to participate will be coming up on [DATE] at [TIME]. The link to join the interview is [HYPERLINK]. Please try to login at least 15 minutes before [TIME] so that the technical support team can assist you, if needed. If you have any questions or find that you are unable to attend, please call [*Insert facility's phone number*] as soon as possible.

Thank you.

[FACILITY INFORMATION]