

Attachment 34_R: No Response – Reminder Letter - 18

Form Approved
OMB No. 0910-0753
Exp. Date 09/30/2019
RIHSC No. 15-101CTP

No Response – Reminder Letter - 18 : FDA Health and Media Study

[CASE ID]

[CHILD'S FIRST NAME OR INITIALS]

[Date]

[Address1]

[Address2]

[City], [State] [Zip]

Dear [CHILD'S FIRST NAME OR INITIALS]:

This is a reminder to complete the fourth follow-up survey for the **FDA Health and Media Study**.

Your participation is important and will contribute to valuable research related to youth and young adult attitudes toward health, health behaviors, and advertisements they may have seen on TV, online, or heard on the radio.

To complete the on-line questionnaire on a personal computer, laptop, phone, or tablet, you must follow all three steps below:

- 1. Open your web browser and type in the study website address: [RUSTEC WEBSITE]**
- 2. Once you have reached the study website, type in the username and password exactly as shown below:**

Username: [Case ID]

Password: [Password]

- 3. Once you've typed in your username and password, you will see instructions for completing this round of the study.**

Those who complete the survey on-line will receive a check for \$20 for completing the survey. If you complete the survey with one of our field interviewers, you will be offered \$20 in cash.

Your help with this round of the study is voluntary. All information that you provide will be kept private to the fullest extent allowable by law and used only for statistical purposes. You or your household will never be identified in any analysis, reports, or publications, and no one will try to sell you anything.

For more information about the study, you can call our project assistance line toll-free at (866) 214-2039, or email us at mediastudy@rti.org. If you have a question about your rights as a study participant, you can call RTI's Office of Research Protection toll-free at (866) 214-2043. Thank you in advance for your help.

Sincerely,

Matthew Farrelly, PhD
RTI International

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Paperwork Reduction Act Statement: The public reporting burden for this collection of information has been estimated to average 5 minutes per response. Send comments regarding this burden estimate or any other aspects of this collection of information, including suggestions for reducing burden to PRASStaff@fda.hhs.gov.