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IRB Authorization Agreement

CDC relying on a non-CDC IRB

This IRB authorization agreement is suitable for documenting a formal agreement between the Centers for Disease Control and Prevention (CDC) and an institutional review board (IRB) on which CDC relies for review of the research activities specified below. This agreement is permitted by human research regulations at 45 CFR 46.114 and 21 CFR 56.114.

1 *Institution or organization providing IRB review (Institution A)*

Name of Institution or Organization A: **NORC at the University of Chicago IRB**
IRB registration #: IRB00000967 IRB registration expiration date: 07/03/2021
Federalwide Assurance (FWA) #, if any: 00000142 FWA expiration date: 07/19/2023

2 *Institution relying on designated IRB (Institution B)*

Centers for Disease Control and Prevention (CDC)
FWA #: FWA00001413 FWA expiration date: 07/05/2022

3 *Scope of authorization agreement*

The officials signing below agree that CDC may rely on the designated IRB at **NORC at the University of Chicago** both for review under 45 CFR part 46 (and 21 CFR parts 50 and 56, if applicable) and for continuing oversight of the involvement of human subjects in the research described below:

	Institution/Organization A: NORC at the University of Chicago	**Institution B: CDC**
Title of research protocol	Social Media for Cancer Prevention	Using Social Media for Recruitment in Cancer Prevention and Control Survey-Based Research
Protocol reference ID	16.06.09	7146
Principal investigator (name, phone, fax, e-mail)		Juan L. Rodriguez, MPH, MS 770-488-3086 (Phone) 770-488-4286 (Fax) fph4@cdc.gov
Primary Contact (name, phone, fax,e-mail)	Michael Stern, PhD 312-357-3891 (Phone) stern-michael@norc.org	Same

Sponsor or funding agency: CDC Award number, if any:

Additional comments:

The review and continuing oversight performed by the designated IRB will meet the human subjects protection requirements of the HHS regulations (and FDA regulations, if applicable) for the protection of human subjects, as well as the requirements of CDC's FWA. The IRB at **NORC at the University of Chicago** will follow written procedures for reporting its findings and actions to appropriate officials at CDC. Relevant minutes of IRB meetings

and related records will be made available to CDC upon request. CDC remains responsible for ensuring compliance with the IRB's determinations and with the terms of CDC's FWA. This document must be kept on file at both institutions and provided to OHRP upon request.

4 Signatures

Institution/Organization A: NORC

Kathleen E Parks 27 August 2018

Signature Date

Kathleen E Parks MBA, MPM
Vice President
1155 East 60th Street
Chicago, IL 60637
312 953-2590 (Phone)
312 759-4004 (Fax)
parks-kathleen@norc.org

Institution B: CDC

Denise M. Marshall 5 Sep 18

Signature Date

Denise M. Marshall, BS
IRB Administrator
1600 Clifton Rd NE MS D-73
Atlanta, GA 30333
404-639-4821 (Phone)
404-639-3249 (Fax)
huma@cdc.gov