



**PERIODIC REVIEW FORM
FOR
RESEARCH INVOLVING HUMAN SUBJECTS**

Please return by: on _____

.....
Instructions: Fill out the form *completely*. Periodic Review cannot be accomplished unless the progress report is completed, a copy of the current consent form is included, and appropriate signatures are obtained. Incomplete forms will be returned.

PRINCIPAL INVESTIGATOR(S): _____

CO-INVESTIGATOR(S): _____

CDPH SPONSOR: _____

DIVISION/PROGRAM: _____

PROJECT TITLE: _____

FUNDING AGENCY OR RESEARCH SPONSOR: _____

PRINCIPAL INVESTIGATOR'S INFORMATION: _____

Mailing Address: _____

E-Mail Address: _____

Telephone Number: _____

Fax Number: _____

.....
 The project was **FIRST APPROVED** on _____

The project was **LAST APPROVED** on _____

.....
I. PROJECT DESCRIPTION

A. Project activity STATUS is (check one of four boxes, as appropriate):

- CONTINUING** with **NO CHANGES** in procedure, risks, or class of human subjects since the last review.
- REVISED.** Minor changes may be indicated on this form. For substantial changes, a new Human Subjects Review Form must be completed, indicating the manner in which the project was revised, and returned with this form. Please complete this form also.
- NEVER INITIATED. WORK WILL NOT BE DONE AT THIS TIME.** Please sign on page three and return this form through the appropriate offices for signature.
- COMPLETED. NO FURTHER CONTACT WITH HUMAN SUBJECTS IS PLANNED.** Please sign on page three and return this form through the appropriate offices for signature.

- B. This project is being conducted at the following SITE(S):
- CDPH CLINICS (please list which sites) _____
 - CDPH FIELD SITES (please list which sites) _____
 - In the field (homes, streets, non-CDPH clinical and/or non-clinical settings) _____
 - Cermak Health Services/Cook County Jail (requires letter from jail co-sponsor) _____
 - Other (specify) _____

II. PROGRESS REPORT

- A. NUMBER OF SUBJECTS STUDIED TO DATE: _____
- NUMBER OF SUBJECTS STUDIED SINCE THE LAST REVIEW: _____
- NUMBER OF SUBJECTS YET TO BE STUDIED: _____

- B. Have any ADVERSE EVENTS been noted since the last review?
- YES NO If YES, how many? _____

Were any of these UNANTICIPATED REACTIONS? ("unanticipated" being defined as not having been anticipated in the protocol nor stated in the consent form)

- YES NO

If you have answered YES to either of the above, please attach an explanation.

- C. Provide a STATEMENT regarding the STATUS of any DRUGS, BIOLOGICS or DEVICES employed in the study.

- D. SUMMARY OF RESULTS to date:

Please attach a copy of your **CURRENT CONSENT FORM**.
Please attach copies of current IRB approval or renewal from collaborating institutions.
Attach copies of any abstracts or publications resulting from this work

Any **ADDITIONAL INFORMATION** that may be useful to the reviewers.

I certify that the approved protocol and approved method for obtaining informed consent have been followed during the period covered by this **PROGRESS REPORT**.

Date

Principal Investigator

III. INSTITUTIONAL ENDORSEMENTS

Your endorsement is requested to assure the Institutional Review Board that your office is aware of the existence and status of this research activity:

Date

CDPH Sponsor

Date

Division Director

.....

Submit to:

Joslyn James
Staff Assistant
Chicago Department of Public Health
333 South State Street
Chicago, Illinois 60604
Email: joslyn.james@cityofchicago.org
(312) 747-8524

[PDF version of this form](#)