Final Progress Report Template

Date (mm/dd/yyyy):

PCORI Project Title:

PI Name (Last, First):

Recipient Organization:

PI Telephone Number:

PI Email:

A. FINAL SCIENTIFIC PROGRESS REPORT

1. Summarize the following: 1) study methods, 2) study protocol, 3) key findings, and 3) interpretations of the relevance of findings to patients, clinicians, and health care systems. You must also attach a complete, final study protocol to this report.

2. Discuss any possible differences in study findings or conclusions among patient subgroups defined by age, gender, race/ethnicity, socio-economic, clinical, or genetic makeup (if studied):

3. Discuss the limitations of study findings in terms of statistical precision, failure to include or study patient subgroups, and possible mismeasurement of key variables:

4. Describe the impact and outcome of the project as it relates to patients:

5. Describe study datasets, including codebooks, meta-data related to the datasets, and documented programming code used for creating the final study population, for creating variables, and for conducting all outcomes analyses:
5. We certify that this project adhered to the PCORI Methodological Standards, as described in the PCORI Methodological Report:

___ Yes  ____ No

If no, please justify:

6a. Notification for Public Acceptance: Did you have any publications or presentations accepted for publication or presentation that you have not reported to PCORI?

___ Yes  ____ No

6b. If yes, list any publications/presentations related to the research. Include publication/presentation title, date published/presented, and where publication/presentation was published or presented. Attach copies of publications and presentations, if relevant.

6c. Notification of Peer Review Rejections: List any scientific article submitted for publication and rejected. Include publication title, date rejected, and where article was submitted.

7. Describe the plan to disseminate the funded research. Include any potential facilitators and/or barriers to dissemination. Describe potential for results to be incorporated into practice.

8. Add any additional findings or comments.
B. FINAL NARRATIVE (NON-TECHNICAL) PROGRESS REPORT

Complete a narrative summary of the report covering the same elements described in A, above, written in language understandable to patients and providers. This report must not include any proprietary information; it will be made public on the PCORI website or through other means. Limit 3 pages.

1. Summarize the following: 1) study methods, 2) study protocol, 3) key findings, and 3) interpretations of the relevance of findings to patients, clinicians, and health care systems:

2. Discuss any possible differences in study findings or conclusions among patient subgroups defined by age, gender, race/ethnicity, socio-economic, clinical, or genetic makeup (if studied):

3. Discuss the limitations of study findings in terms of statistical precision, failure to include or study patient subgroups, and possible mismeasurement of key variables:

4. Describe the impact and outcome of the project as it relates to patients:
5. Describe the plan to disseminate the funded research. Include any potential facilitators and/or barriers to dissemination. Describe potential for results to be incorporated into practice.

6. Add any additional findings or comments.
CERTIFICATION: We, the undersigned, certify that the information submitted is accurate and complete to the best of our knowledge and accept the terms and conditions of PCORI as described in the PCORI Contract. Please note: all required signatures must be submitted.

SIGNATURES: Principal Investigator ____________________________________________

Administrative Official _____________________________________________________