Study Progress Report
Relying Site Principal Investigator

CHOP IRB Study #

Protocol Title:

Expiration Date:

Relying Site Principal Investigator:
Relying Institution:

1. Select Report Type: Continuing Review Report ◯
   Closure Report ◯
2. Enrollment Status: Enrolling Subjects ◯
   Enrollment Closed ◯
3. # Subjects Enrolled since the last IRB (Initial or Continuing) review:
4. How many subjects at your site withdrew from the study or were withdrawn by the investigator*?

   Reasons for the withdrawal(s):
5. Have subjects at your site registered any complaints about the research or study personnel since the last CHOP IRB approval?
6. Provide (or attach) a summary of any noteworthy adverse events or other study-related issues that occurred at your site and did not meet the criteria for prompt reporting to the CHOP IRB (e.g. Unanticipated Problems involving risk to subjects or others or non-compliance).
7. Provide a summary and explanation of any Adverse Events or Unanticipated Problems that have occurred at your site with an unexpected frequency or severity, that have not been previously reported since the last CHOP IRB approval.
8. Have there been any material changes to the Relying Site Survey, which were not previously reported to the CHOP IRB? If so, briefly outline the nature of these changes.

Confirm the following:

- The Relying Site Survey, and all documents submitted to the CHOP IRB are current

and accurate.

- All Unanticipated Problems, that met reporting criteria, were submitted to the CHOP IRB as Reportable Events. (CHOP IRB Policies are available at https://irb.research.chop.edu/policies)

Relying Site Principal Investigator Signature: