The 5th Institutional Review Board Meeting (FY2018) (Expedited review)

Date: August 21, 2018 (Tuesday) 14:00-16:40

Place: Auditorium (Hiroshima), 3rd fl. meeting room (Nagasaki) (teleconference)

Participants: Dr. Kodama and Ms. Shinohara (co-chairpersons), Dr. Yamada, Dr. Satoh, Dr. Sugiyama, Dr. Hida, Ms. Dodo and Ms. Ogawa (members)

＜Review concerning human subject research＞

	Department	Research project title	Review results	Department
1	Clinical Studies	Revision of RP1-17 “Detection of the onset of hematological malignancy among atomic bomb survivors” (Imaizumi *et al.*)	Approved	● The RP will be revised in the following manner: 1) Revise description of data storage/management. 2) Detail new researchers in revised chart 2. 3) Check how “genome data” and “genome information” are defined legally and in the ethical guidelines, and revise the relevant descriptions accordingly. ● Revise the description of “Safeguarding personal information” on the consent explanatory note.
2	Clinical Studies	CR149 “Research protocol for collection of radiation dose data from basic health surveys of Fukushima residents” (Addendum to RP6-15 “Epidemiological Study of Health Effects in Fukushima Emergency Workers”) (Okubo *et al.*)	Approved	● The RP will be revised in the following manner: 1) Describe the participants in the RP properly. 2) Ensure the description of “informed consent” matches the expressions on the consent form. ● Revise the expressions on the “Request for provision of your basic health survey results” in an easily understandable manner for participants. ● The Consent/response form should correctly detail the name of the research project for which the information will be used.

*With regard to the aforementioned issues, IRB confirmed that proper revisions were made.