AAHRPP INFORMATION ITEM

July 1, 2019

Association for the Accreditation of Human Research Protection Programs, Inc.

2301 M Street, NW

Suite 500

Washington D.C. 20037

[accredit@aahrpp.org](mailto:accredit@aahrpp.org)

Dear AAHRPP:

On *<Receipt Date>* the IRB was notified of the following information item(s): *<delete all that do not apply>*

* Negative actions by a government oversight office, including, but not limited to, OHRP Determination Letters, FDA Warning Letters, FDA 483 Inspection Reports with official action indicated, FDA Restrictions Placed on IRBs or Investigators, and corresponding compliance actions taken under non-US authorities related to human research protections.
* Litigation, arbitration, or settlements initiated related to human research protections.
* Press coverage (including but not limited to radio, TV, newspaper, online publications) of a negative nature regarding the Organization’s HRPP.

This information is in regard to:

* *<Insert description. Delete this section if no information is required.>*

Please let us know if you need additional information at this time.

Sincerely,

IRB Manager