cbj 12.07.22

EXPEDITED REVIEW PETITION FORM

for NUNM's Institutional Review Board (IRB)

Expedited Status Research

Expedited review is a procedure through which certain kinds of research may be reviewed and approved without convening a meeting of the entire IRB. The term "expedited" does **not** mean a review is quicker or conducted with less rigor. It means fewer reviewers are required for approval. In general, research may be considered for expedited review if it involves no more than minimal risk, does not include intentional deception, does not employ vulnerable populations or sensitive topics, and includes appropriate consent procedures.

Please note that **all of the rights and protection afforded to human subjects in research are required in expedited status cases.** Investigators engaged in human subject research that qualifies for expedited status **must still complete an IRQ, prepare an informed consent statement (as appropriate for the methods), copies of any/all data collection instruments (e.g., survey, standardized interview guide, data abstraction forms, etc.), copies of any/all materials that will be viewed or received by participants, and provide a detailed description below of the project aims, data collection methods, and analytical plan for any data collected.**

Please email this completed form to the IRB Liaison at IRB@nunm.edu with all required documents for IRB review.

Study Title:

Short Title or Acronym:
(limit to 54 characters)

PI Telephone Number: **PI Email:**

Submission with petition for EXPEDITED review - this project is (check the relevant option):

☐ A clinical study of drugs for which an investigational new drug application is not required.

☐ Research on medical devices for which an investigational device exemption application is not required; or the medical device is cleared/approved for marketing and the medical device is being used in accordance with its cleared/approved labeling.

☐ Research involving collection of blood samples by finger stick, heel stick, ear stick, or venipuncture per the Office for Human Research Protections expedited review guidelines.

☐ Research involving prospective collection of biological specimens for research purposes by noninvasive means.

☐ Research involving collection of data on subjects 18 years of age or older through noninvasive procedures routinely employed in clinical practice.

☐ Research involving materials that have been collected, or will be collected solely for non-research purposes.

☐ Research involving collection of data from voice, video, digital, or image recordings made for research purposes.

☐ Research on individual or group characteristics or behavior; or research employing survey, interview, oral history, focus group, program evaluation, human factors evaluation, or quality assurance methodologies.

Please detail the overall study methodology below (e.g., survey, retrospective medical chart review, demonstration project of service program, etc.). Provide more detail than the checklist above allowing reviewers to fully understand what you intend to do in your proposed research.

Please provide the Aims/Objectives of you proposed research:

Provide details on your analytical plan for any/all data collected:

☐ The study team confirms copies of all an informed consent statement (as appropriate for the methods), copies of any/all data collection instruments (e.g., survey, standardized interview guide, data abstraction forms, etc.), and copies of any/all materials that will be viewed or received by participants are included with this submission.

PI Signature Date

This section is for the IRB Chair only

Review Date: **IRB Number:**

IRB Expedited Review Approved (Please see comments on original documents.)

If an expedited review is approved, file this form along with all proposal documents where they can be easily accessed and identified.

IRB Expedited Review Not Approved (Please see comments on original documents.)

If the expedited review is not approved, the IRB Liaison will contact the PI to inform them that the study has not been approved for expedited review and will be reviewed by the full IRB.

IRB Chairperson Signature Date