

**ANNUAL REPORT / RENEWAL - INVESTIGATIONAL HUMAN-USE APPLICATION
APPROVED TO USE RADIOACTIVE MATERIALS (RAM) AND/OR RADIOGRAPHIC MACHINES**

(Check boxes wherever applicable)

RAM only

RADIOGRAPHIC MACHINES only

RAM & MACHINES

1) TITLE OF THE RESEARCH PROJECT:

2) PRINCIPAL INVESTIGATOR (PI) Information (Please Print Clearly) :

Last Name: _____ First Name: _____ M.I. _____
Title: _____ Department: _____
Office Location: _____ E-mail: _____
Office Phone # _____ Emergency Phone # _____ Lab Phone # _____

3) Number of Human Research Subjects Studied since last renewal (12/1/20 to 11/30/20) :

4) Cumulative Number of Human Research Subjects Studied over the Lifetime of this Protocol :

5) Since the last review, have **any unanticipated problems involving risks to subjects or others and/or serious adverse events occurred** that have not been previously reported?

Yes No *If yes, please attach a copy of detailed explanation of each event.*

6) Since the last review, were **the radiation doses within the specifications of the approved protocol for all subjects?**

Yes No *If No, please provide detailed calculation of absorbed doses and effective doses for each research subject that exceeded the dose estimates in the approved protocol. Dosimetry for radiopharmaceuticals should include absorbed dose and equivalent dose to whole body, active blood-forming organs, lens of the eye, gonads, and other organ doses. Also provide effective dose.*

7) Was a claim of confidentiality made?

Yes No

Note: *Contents of these reports will be made available for public disclosure unless confidentiality is requested by the investigator and it is adequately shown by the investigator that the report constitutes a trade secret or confidential commercial information.*

8) Signature below affirms that the Principal Investigator will comply with the requirements set forth by the Radiation Safety Committee for the use of radionuclides and machine sources of ionizing radiation. Furthermore, the Principal Investigator shall notify the Human Use Subcommittee of the Radiation Safety Committee immediately of any adverse reactions that may be attributable to radiopharmaceuticals or machine source radiation administered to their research subjects.

In case of prolonged absence or termination, the Principal Investigator must notify the REHS-Radiation Safety Officer by telephone 973-972-5305 or E-Mail: Prasad.Neti@rutgers.edu.

Signature

Date