RDRC#: 185

RDRC Protocol number:

Project Title:

Principal Investigator:

Reporting period:

Report Date:

N.B. Do not provide any protected health information. If you need to identify subjects, provide a protocol tracking number (e.g. subject number 1 for the first subject enrolled). This is only needed for consistency of identification throughout the course of the protocol.

Please provide the following information.

• A summary of all patients recruited into each protocol and subgroups, if any, for the reporting period:

• The number of extra doses of radiopharmaceuticals administered due to technical failures as part of proposed protocol contingencies:

• Issues related to radiochemistry production and quality assurance:

• Adverse events. An Adverse Event is defined as any event associated with the radiotracer that are not clearly defined in the protocol as expected side effect of other drugs, which may be used in the protocol: