Form B

Institutional Review Board Le Moyne College

APPLICATION FOR EXPEDITED REVIEW

Title of Project:
Name of Primary Investigator(s)/Researcher(s):
Type of Investigator(s):
Faculty or staff of Le Moyne College
Student of Le Moyne College
Individuals other than faculty, staff, or students of Le Moyne College
(Le Moyne researchers must identify all outside investigators / collaborators and explain their role on the Research Outline.)
Name of Student Research / Lab Assistant(s):
Name of Student Research / Lab Assistant(s): (On the Research Outline, explain the role of student research / lab assistants who will assist in data
collection or analysis but will not be otherwise involved in preparing the final products.)
CITI Completion Certificates must be submitted for all primary investigators, researchers / collaborators
student research / lab assistants, and faculty sponsors.
Primary Investigator's Contact Information:
Address
City, State & Zip Code
Phone number
E-mail address
For Student Researchers only:
Degree program (e.g., BA, MS, EdD, DNP):
Program(s) of Study:
Name of faculty sponsor:
Email address of faculty sponsor:
For External Researchers only:
Name of Le Moyne sponsor:
Email address of Le Moyne sponsor:
Date Application Submitted:
Proposed Date to Commence Data Collection: (Please allow a minimum of 6-8 weeks between the application submission date and the proposed date to
commence data collection.)

Natu	re of Activity (check all that apply):		
	Faculty research project Project to be submitted for extramural funding from a federal agency (e.g., NSF, Agency: Grant Submission Deadline:		_
	 Pilot Study for future research project Undergraduate thesis / capstone project (including Departmental / Integral Hono Master's thesis / capstone project Doctoral dissertation 	ors)	
	Course project (not thesis or capstone related). Course number & title: Staff (non-faculty) research project Other (please explain):		
Abst	ract of Project (maximum 150 words):		
	se answer the following questions with regard to the proposed research activity. (A mative response to any of these might necessitate a Form C with Full Board review.)		
Does	the research involve:	YES	NO
a.	drugs or other controlled substances?		
b.	participants taking internally or having externally applied any substance(s)?		
c.	removing any fluids (e.g., blood) or tissue from participants?		
d.	participants experiencing stress (physiological or psychological) above a level that would be associated with normal everyday activities?	: 	
e.	misleading (deceiving) participants about any aspect of the research?		
f.	participants who would be judged to have limited freedom of consent (e.g. minors or individuals with impaired decision-making ability)?		
g.	any procedures or activities that might place the participants at more than minimal risk (psychological, physical, or social)?		
h.	sensitive aspects of the person's own behavior, such as illegal conduct, drug use, sexual behavior, or alcohol use?		
i.	activity or data collection whereby the disclosure of participants' responses outside the research could place the participants at risk of criminal or civil liability or be damaging to the participants' financial standing, employability, educational advancement, or reputation.		

Moderate exercise by healthy volunteers. The study of existing data, documents, records, pathological specimens, or diagnostic specimens, if the individuals from whom the data collected are identifiable. Research on individual or group behavior or characteristics of individuals, such as studies of perception, cognition, game theory, or test development, where the investigator does not manipulate participants' behavior and the research will not involve stress to participants. Collection of: hair and nail clippings, in a non-disfiguring manner; deciduous teeth; and permanent teeth if patient care indicates a need for extraction. Collection of excreta and external secretions including sweat, uncannulated saliva, placenta removed at delivery, and amniotic fluid at the time of rupture of the membrane prior to or during labor. Recording of data collected from participants 18 years of age or older in the course of noninvasive procedures routinely employed by professionally certified/licensed individuals in the clinical practice of medicine, psychology and social work. This includes the use of physical practice sensors that are applied either to the surface of the body or at a distance and do not involve input of matter or significant amounts or energy into the research participant or an invasion of the participant's privacy. It also includes such procedures as weighing, testing sensory acuity, electrocardiography, electroencephalography, thermography, detection of naturally occurring radioactivity, diagnostic echography, and electroretinography. It does not include exposure to electromagnetic radiation outside the visible range (e.g. x-rays, microwaves.) Collection of blood samples by venipuncture, in amounts not exceeding 450 milliliters in an eight-week period and no more often than two times per week, from participants 18 years of age or older and who are in good health and not pregnant.

Under which of the following categories are you applying for EXPEDITED REVIEW? (check one)

Certification:

accordance with accepted prophylactic techniques.

1. I am familiar with the policies and procedures of Le Moyne College regarding human research participants. I subscribe to the standards described in the document, *Institutional Review Board Policies and Procedures*.

Collection of both supra- and subgingival dental plaque and calculus, provided the procedure is

not more invasive than routine prophylactic scaling of the teeth and the process is accomplished in

- 2. I am familiar with the published guidelines for the ethical treatment of human research participants associated with my particular field of inquiry (e.g., as published by the American Psychological Association, American Sociological Association.)
- 3. I am familiar with and will adhere to any official policies in my department concerning research

with human research participants.

- 4. I understand that upon consideration of the nature of my project, the IRB may request a Form C application for full Board review of my research at their discretion.
- 5. If changes in procedures involving research participants become necessary, I will submit these changes for review before initiating the changes.

SIGNATURE Investigator/Researcher	DATE	_
SIGNATURE	DATE	_
All student-led research application have a faculty / college sponsor wh	ons and all applicants from outside Le Mo hose signature is here affixed:	yne College must
SIGNATURE	DATE	_
Additional signatures (as needed):		
SIGNATUREInvestigator/Researcher	DATEOther:	_
SIGNATURE Investigator/Researcher	DATE Research / Lab AssistantOther:	_

Updated June 2023