ETHICS FORM

IDENTIFICATION
Title of the study or project
Contextualization of the study or project (or Scope of the research (e.g. PhD, research project)
Name of the proponent/person responsible for the study
Name of the supervisor/scientific officer of the study (if different from the previous one)
Contact for questions or information (e.g. email, phone)
PREVIOUS SUBMISSIONS to Ethics Committees
Has this paradigm ever been submitted to an Ethics Committee?
Has this paradigm been previously approved by this Ethics Committee?
Are there any changes to what was previously approved? Describe the changes.
DESCRIPTION (Can be replaced by a thesis project/project funded by FCT). DESCRIBE, IN ONE PARAGRAPH:

Objective and framework of the research to be carried out:
Methodological design of the study (Phases and Procedures of manipulation and measurement):
Participants: characterization, selection/exclusion criteria, form of recruitment and/or referral, etc.
Procedures: timetable, phases, instructions and procedures for data collection, etc. (specify only those aspects that may have ethical implications for the type of sample concerned)
Instruments: identification (e.g. questionnaires, observations, document analysis, physiological measurements) and their framing
Information given to participants and special care in case of non-relevant placement (NR)
Intrusive study? Special care? Reparation of any damage or inequality by manipulation/intervention? (in case of not relevant put NR)

Informed consent?
Voluntary nature of participation?
Does the investigation involve minors? If so, what consent for the legal representatives? If the minor has the capacity to understand and manifest his will, his assent is also required.
How is confidentiality guaranteed in the treatment and dissemination of results? How to preserve the collected information (recordings, biological data, etc.) for the future?
END-OF-STUDY PROCEDURES
Will participants and communities be informed of the outcome of the study? Why means? (e.g. overall results returned to the participant or focus group; individual results to the participant... If so, how:
Is it intended to be disseminated to the scientific community?
CONFLICTS OF INTEREST
Does anyone involved in the design and/or development of the research have particular interests that are related to the elements analysed in the study (e.g. patents, trademarks, copyrights)?

DOCUMENTS TO BE ATTACHED

Attach the required documents illustrating all materials, methods, and procedures (e.g., informed consent letters, questionnaires, scales, interview guides, or physiological measurement protocols).