Attachments:

- Appendix 8_Informed Consent Form_ Focus Group Participation.pdf
- Appendix 7_The Respondent Informed Consent Model (e.g. Consent Form for Teacher Interview Participation).pdf
- Appendix 9_The Informed Consent for participants to U-report survey.pdf
- Appendix 6_Child-Friendly Assent Form (Ages 11 14) - Focus Group Participation.pdf
- Appendix 5_Parents guardians of adolescents no objection form.pdf
- Ethical Review Approval.pdf

HML IRB
RESEARCH & ETHICS

Ethical Review Approval

To: Rachid AMRI, MA

From: HML IRB

Subject: Study #3010

Date: 17-Jul-2025

Dear Rachid AMRI, MA,

The protocol **Evaluation of the project "Accompaniment of adolescents and young people in their transition from adolescence to adulthood, 3010** was assessed through a research ethics review by Health Media Lab Institutional Review Board. This study's human subjects' protection protocols, as stated in the materials submitted, received research ethics review approval on 17-Jul-2025.

You may rely on this IRB for review and continuing ethical oversight of this study. You and your project staff remain responsible for ensuring compliance with the IRB's determinations. Those responsibilities include, but are not limited to: 1) ensuring prompt reporting to HML IRB of proposed changes in this study's design, subject risks, informed consent, or other human protection protocols and providing copies of any revised materials; 2) conducting the research activity in accordance with the terms of the IRB approval until any proposed changes have been reviewed and approved by the IRB, except when necessary to mitigate hazards to subjects; 3) promptly reporting any unanticipated problems involving risks to subjects or others in the course of this study; and 4) notifying the IRB when your study is complete.

The approval of your study is valid through 16-Jul-2026, by which time you must submit a continuing review report either closing the study or requesting permission to continue for another year. Please submit your report by so that the IRB has time to review and approve your report prior to the expiration date. For instructions on how to manage an approved study refer to: How to Manage an Approved Study.

Health Media Lab IRB is authorized by the U.S. Department of Health and Human Services, Office of Human Research Protections (IRB #00001211, IORG #0000850), and has DHHS Federal-Wide Assurance approval (FWA #00001102).

If you have any questions, please contact us at admin@hmlirb.com.

Sincerely,

D. Michael Anderson PhD, MPH
IRB Chair & Human Research Protections Director
dma@hmlirb.com