

Office of Research INSTITUTIONAL REVIEW BOARD

MEMORANDUM

To: Scott Rhodes, Ph.D.

PHS-Social Sciences

From: Deborah Wesley, Assistant Director, Institutional Review Board

Date: 9/11/2019

Subject: Human Protocol: IRB00040441

HIV Prevention among Latina transgender women who have sex with men: Evaluation of

a locally developed intervention

Amendment 28 for IRB Study #IRB00040441 (Personnel Change Request)

Study Documents:

Protocol Version: ChiCAS Protocol (08 22 19).docx; Informed Consent Version: Appendix C1_English Consent Form.doc.docx (approved), Appendix C2 Spanish Consent Form.docx (approved); Advertisements: Appendix H1 English Informational Flyers.pdf, Appendix H2 Spanish Informational Flyers.pdf; Other Documents: Appendix B1_English Participant Screening Form.docx, Appendix B2_Spanish Participant Screening Form.docx, Appendix D1_English Questionnaire.docx, Appendix D2_Spanish Questionnaire.docx, Appendix E_Spanish-language ChiCAS Curriculum Manual.docx, Appendix F_Implementation Observer's Guide.docx, Appendix G_Participant Attendance Log.docx, Appendix I_Medical Chart Abstraction Form.docx, Appendix J_Data Sharing and Use Agreement.docx, Appendix K1 English Qualitative In-Depth Interview.docx, Appendix K2 Spanish Qualitative In-Depth Interview.doc, Appendix L1 English Stay In Touch Cards.docx, Appendix L2 Spanish Stay In touch Cards.docx, Appendix M1 English Appointment Card.docx, Appendix M2 Spanish Appointment Card.docx, Appendix N_Individual Investigator (Trent) IRB Agreement.pdf, Appendix O_UNC Greensboro IRB Agreement.pdf, Appendix P1 English script for DVD1 (HIV impacts).docx, Appendix P2_Spanish script for DVD1 (HIV impacts).docx, Appendix Q1_English outline for DVD2 (PrEP etc.).docx, Appendix Q2_Spanish script for DVD2 (PrEP etc.).docx, Appendix R_Hormone Therapy PPT curriculum (Sp).pptx, Appendix S HIV and STDs PPT curriculum (Sp).pdf, Appendix T PrEP PPT curriculum (Sp).pptx

The amendments listed below have been approved in accordance with HHS regulations for the protection of human research subjects that provides for the expedited review and approval of minor changes in previously approved research [45 CFR 46.110(b)(2)]. This action of the Board does not extend the term of approval for this protocol.

The amendment includes the following:

Adding Lisa Lewis as study coordinator.

The Wake Forest School of Medicine IRB is duly constituted, has written procedures for initial and continuing review of clinical trials; prepares written minutes of convened meetings, and retains records pertaining to the review and

approval process; all in compliance with requirements of FDA regulations 21 CFR Parts 50 and 56, HHS regulations 45 CFR 46, and International Conference on Harmonisation (ICH) E6, Good Clinical Practice (GCP), as applicable. WFSM IRB is registered with OHRP/FDA; our IRB registration numbers are IRB00000212, IRB00002432, IRB00002433, IRB00002434, IRB00008495.

WFSM IRB has been continually fully accredited by the Association for the Accreditation of Human Research Protection Programs (AAHRPP) since 2011.

