Complete this form when a local IRB has jurisdiction over this research site.

You must submit a typed version of this form (except for the signature block) to prevent errors and delays due to legibility problems.

Blank & incomplete answers will result in delayed reviews

If you have questions about the use of this form, please contact WCG IRB at *855-818-2289* or email *clientcare@wcgirb.com*

Relying Organization:

Name:	San Diego State University IRB
FWA #:	00003782 (if organization has an FWA)

Contact Information for Relying Organization:

Name:	Anne Dodge-Schwanz		
Title:	Lead IRB Analyst		
Address 1:	5500 Campanile Drive		
Address 2:	MC 1933		
City:	San Diego	State/Province:	CA
Zip/Postal Code:	92182	Country:	United States
Phone:	619-594-6622	Email:	adodge@sdsu.edu

☐ **This agreement covers all human subjects research conducted by this site** (otherwise complete the "Research Protocol" and "Principal Investigator" sections below)

Research Protocol:

Protocol #:	None
Protocol Title:	Understanding HIV/STD Risk and Enhancing PrEP Implementation Messaging in a Diverse Community-Based Sample of Gay, Bisexual, and Other Men Who Have Sex with Men in a Transformational Era (MIC-DROP)
Sponsor Name:	Centers for Disease Control and Prevention
IRB Tracking #:	20225896 (if known)

Principal Investigator:

Name:	Patrick Sullivan, DVM, Ph.D. (Emory), Keith Horvath, Ph.D. (San Diego State University)

The Relying Organization agrees to waive jurisdiction for the IRB review and continuing oversight of the Research Protocol to WCG IRB (IRB registration number IRB00000533) ("Designated IRB") as allowed under federal regulations. The Principal Investigator is authorized to conduct the Research Protocol at the Relying Organization.

The review performed by the Designated IRB will meet the human subject protection program requirements of applicable regulatory agencies. The Designated IRB will follow its written procedures for the review and oversight of the Research Protocol. The Designated IRB will follow its written procedures for reporting findings and actions, subject complaints, unanticipated problems involving risks to subjects or others, serious or continuing noncompliance, and suspension or termination to appropriate institutional officials at the Relying Organization. Relevant minutes of IRB meetings will be made available to the Relying Organization upon request. Relying Organization remains responsible for ensuring compliance with the Designated IRB's determinations and with the terms of any

agreements between the Relying Organization and applicable regulatory agencies. This document must be kept on file by both parties and provided to applicable regulatory agencies upon request.

For Designated IRB

DocuSigned by:
Sheri Spivey
Signature

Sheri Spivey
Printed Name

Regulatory Chair Director
Title

20-Mar-2023 | 11:36 AM EDT
Date

For Relying Organization

Natalie Gude
Signature

Natalie Gude, Ph.D.
Printed Name

Assistant Director for Research Support Services
Title

14 March 2023
Date