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:	University of Chicago BSD-IRB
FWA #:	00005565 (if organization has an FWA)

Contact Information for Relying Organization:

Name:	Jeremy LaVigne, MA, CIP		
Title:	IRB Reliance Manager		
Address 1:	University of Chicago		
Address 2:	5841 South Maryland		
City:	Chicago	State/Province:	IL
Zip/Postal Code:	60637	Country:	USA
Phone:	(773) 834-8262	Email:	jlavigne@bsd.uchicago.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:	John Schneider, MD

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:
Bridget Brave

Signature

Signer Name: Bridget Brave
Signing Reason: I have reviewed this document
Signing Time: 18-Aug-2023 | 11:00:58 AM EDT
B411625B7E52430E874F4E13BFF1865C

Printed Name

Director, Expedited Review

Title

18-Aug-2023 | 11:01:01 AM EDT

Date

For Relying Organization

Jennifer A Ponting

Signature

Jennifer A Ponting

Printed Name

Associate Vice President for Research, Director of University Research Administration

Title

8/10/23

Date