



Purpose of Form: *If not using the SMART IRB Online Reliance System to coordinate and document study-specific reliance arrangements, institutions may use this template to document the Reviewing IRB and Relying Institutions for a study.*

Acknowledgement of Site Agreement to Cede IRB Review and Reviewing IRB to Provide Oversight

This form documents that:

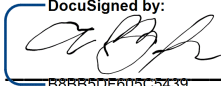
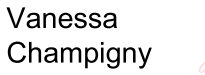
- 1) The Columbia University Medical Center (CUMC) IRB will serve as the Reviewing IRB for **University of Alabama at Birmingham** for the study noted below;
and
- 2) **University of Alabama at Birmingham** has agreed to cede IRB review to the CUMC IRB for the study noted below.

Study Title:	mChoice: Improving PrEP Uptake and Adherence among Minority MSM through Tailored Provider Training and Adherence Assistance in Two High Priority Settings
CUMC Principal Investigator:	Rebecca Schnall, PhD
CUMC RASCAL Study Number:	AAAT8812
Sponsor or Funding Agency (if any):	Centers for Disease Control (CDC)
Relying Institution Principal Investigator:	Mirjam-Colette Kempf, PhD, MPH
Relying Institution Study Number (if any):	IRB-300008573

IRB review will be ceded under the SMART IRB Master Common Reciprocal Institutional Review Board Authorization Agreement Version 1.

Questions about the IRB review process or study status should be directed to Tasha Isles Smith, MA, MPH, at ts2257@cumc.columbia.edu

Acknowledged by:

CUMC IRB Designee Signature:  <small>B8BB5DF605C5439...</small> Print Name: Andrew B. Lassman, MD Title: Associate Dean of Clinical Research Compliance	Date: 2/18/2023 12:26:29 AM GMT
University of Alabama at Birmingham Designee Signature:  <small>Digitally signed by Vanessa Champigny Date: 2023.02.02 16:08:08 -06'00'</small> Print Name: Vanessa M. Champigny, MBA, CIP Title: Assistant Director, UAB Office of the IRB	Date: 02/02/2023