

Institutional Review Board 5 i h cf]nUf]cb'5 [fYYa YbhForm

Omaha, Nebraska 68114 (402) 354-4000	This form is ¦	``å^åÁqíÁ&^å^Áqíç^¦∙ãt@Áofastudyq[Ása)Á^¢ơ^¦}æ‡ÁÜÖÓ.
	RTY A	
Institution/Organization*		
IRB Registration #*		
IRB FWA #*		
PA Institution/Organization*	RTY B	
IRB Registration #*		
IRB FWA #*		
Select One*	•	approval and will oversee all Study conduct.
TITLE of Clinical Research Study*		
Protocol Number*		
Principal Investigator*		
Sponsor*		
Provide Additional Details for Transfer Below:		
The execution of the document confirms Party ma review performed will meet the Human Subject Protection	requirements of OHRP approved F	WA. IRBs will follow written procedures for reporting their
findings and actions to appropriate officials. Relevant mini- ensuring compliance with the determinations and with the provided upon request.	utes of IRB meetings will be made a TERMS of OHRP-approved FWA.	available upon request. Both parties remain responsible for This document will be kept on file by both parties and
PARTY A Authorization*	Print Full Name	
	Sign	Date
PARTY B Authorization*	Print Full Name	
	Sign	Date

* required field Version 1.1 (21Jan2025) Page 1 of 1