Catawba Valley Med Cnt Institutional Review Board IRB-Approved Research Study Withdrawal Form

The Institutional Review Board must be informed in the event a study that has been approved is not conducted. It is the Primary Investigators (PI's) responsibility to inform the IRB so that the study can be withdrawn from IRB monitoring. After receipt and review of a Study Withdrawal form, the IRB will notify the PI that the Board has withdrawn the study from IRB monitoring.

<u>Instructions</u>: Complete and submit this form upon deciding not to conduct the approved study. The Study IRB Number, Approval Expiration Date and Approval Type are contained in the original IRB approval letter received. Submit this form to <u>irb@cvmc.us</u>.

	Date:	
Study Title:		
Primary Investigator (PI) Name:		
PI Email:	PI Phone:	
IRB Approval Expiration Date:		
Original Approval:	Full Board Review	Expedited Review
Withdrawal Request Rational (choose one of the f	following)	
withdrawai request rational (choose one of the f	oliowing)	
1. Research determined not to be feasible at this	time	
 No research activities were initiated 		
2. Primary investigator unable to conduct the stud	dy	
No research activities were initiated		
OTE: If ANY research activity was begun, submit a study	closure form not this study w	ithdrawal form.
, ,		
Electronic Signature: Disclaimer		
By signing your name electronically below, you are agreeing to manual signature on this Study Withdrawal Form.	hat your electronic signature is th	ne legal equivalent of your
Investigator Signature:	Date <u>:</u>	
IRB Us CATAWBA VALLEY MED CNT INSTITUTIONAL REVIEW BOARD SIGNATURE:	se Only	
Date of Study Withdrawn from Monitoring:		

irb_study_withdrawal_from

Created: 2017-10; Revised: 2023-03