Steps to completing the ROUTE SHEET: Save blank ROUTE SHEET to your desktop; Complete the required information; Electronically sign and save the completed form; Last step, forward to respective Parties for their approvals. Thank you!





CLINICAL RESEARCH INSTITUTE Phone: (806) 743-4222 • Suite BA-101 • Mail Stop 8183

THIS FORM MUST BE COMPLETED, SIGNED AND SUBMITTED WITH ALL NEW PROJECTS

Scan and submit to: <u>clinicalresearch@TTUHSC.EDU</u> or mail original to STOP 8183

-			
Principal Investigator	Campus/School	Department	Room/STOP
Phone	Fax	Email Address	
Sub-Investigator	Campus/School	Department	Room/STOP
Sub-Investigator	Campus/School	Department	Room/STOP
Sub-Investigator	Campus/School	Department	Room/STOP
If more than three Sub-Investigators nle	ease use second sheet		

in more than three Sub-investigators, please use second she

Study Title (max 150 characters):

Where will the human subjects/charts be recruited from?				
How will the human subjects/charts be identified?				
Estimated study start & completion dates:				
List study expenses (budget) and how they will be funded: Attach budget details on separate sheet, if needed				
Assistance requested from Institute: (check all that apply)				
IRB Work Data Collection/Chart Reviews	Recruitment/Consenting			
	Specimen Collection			
Statistical Analysis List of documents needed	Abstract/Poster/Manuscript Preparation			
Study Design Complete required form	Other			
Sample Size Complete required form				
Would you be interested in having a student(s) involved in this study? Y N				
Would you be interested in having a resident(s) involved in this	s study? Y N			

Please note, in order to provide adequate support for your study, the Clinical Research Institute (CRI) requires you to submit a Semi-Annual Report Form to the Institute. This report form <u>must</u> be completed every June & December. The principal investigator will receive an email and a reminder before this report is due.

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PUBLICATION

It is important to remember that some sort of publication ensues from your research.

Please note, using the CRI resources requires that you INVOLVE the CRI in the preparation of any presentation, abstract, or publication resulting from this study.

Further, please include an acknowledgement that, "This study was supported in part by the TTUHSC CLINICAL RESEARCH INSTITUTE."

Should you feel that the Director(s), coordinator(s), medical student, and/or resident involved with your study has made a significant intellectual contribution, it is recommended you consider including them as a co-author on any presentation/publication.

INVESTIGATOR: By signing below, I certify that (1) the information submitted within this form is complete and accurate to the best of my knowledge, (2) I accept the responsibility for the scientific conduct of the project, and (3) I have completed the TTUHSC required training for clinical investigators.		DEPARTMENT CHAIR or AD of Research: I have reviewed the protocol and find it consistent with TTUHSC and department policies and objectives. The Investigator has the skills and the department has the available resources (space, equipment, personnel, and funding if applicable) to support this program (there are no costs for the services of the Institute).		
Investigator Signature		Department Chair or Associate Dean of Research Signature		
Print Name	Date Signed	Print Name	Date Signed	
		Regional Dean Signature		
		Print Name	Date Signed	

** PLEASE INCLUDE A COPY OF YOUR PROTOCOL AND DATA COLLECTION SHEET WHEN SUBMITTING THIS FORM **

Investigator Required Training:

- <u>CITI Training</u>
- Financial Disclosure
- <u>iRIS User Account</u>

CRI Office Use Only		
Date Rec'd:		
CRI #:		
IRB App'd:		
IRB#:		