

OFFICE USE ONLY
CRC Project #
Date Rec'd

Clinical Research Center Request for Services Form

Protocol Title:			
Study Short Title:			
Sponsor / Funding Source:			
Sponsor Protocol Number:			
IRB # (if available):		IRB Expiration Date:	
OCR# (if available)			
Principle Investigator:			
PI Department:			
PI Email Address:			
PI Phone Number:			
Co-Investigators:			
Project MD:			
Primary Study Coordinator:			
Study Coordinator Email:			
Study Coordinator Phone:			
and will be attached at the botto	needed from CRC and click the 'su om of this form. he most recent version of this for		
PLEASE MAKE SUI	RE REQUESTED DOCUMENTS ARE A	ATTACHED OR YOUR REQUEST MA	Y BE DELAYED
UF CLINICAL RESEARCH CENTE	R SERVICES NEEDED		
1. What is the participan	t population? (check all that app	oly)	
□ Children	☐ Adults ☐ Elderly		
2. What is the Age Range	of Participants?	years of age	
3. How many participant	s do you plan to enroll?		
4. What is the projected	start date of the study?		
5. What is the anticipate	d duration of the study?	Months	
6. What type of visits wil	l be needed? (check all that app	oly)	
☐ Outpatient vis	its during regular CRC hours (7:	00 am – 5:00pm)	
·	its during regular CRC hours (7:0 rs (5:00pm – 12:00am)	00 am – 5:00pm)	

	Indicate nursing services needed for extended and overnight visits:				
		Scatter-bed services (CRC will provide research nursing coverage for participants in Shands Hospita			
		other facility on campus between 7:00am – 5:0	00pm)		
	Please indicate the services the scatter-bed CRC nurse will provide:				
7.	Do yo	u need a room with: (check all that apply)			
		Hospital Bed Chair Str	retcher		
8.	NUR	SING SERVICES – Please indicate the nursing serv	vices requeste	d: (check all that apply)	
		Vital Signs		BMR	
		Height/Weight		BodPod	
		Blood Draw		Finger Stick	
		Specimen Collection		EKG	
		IV Catheter Placement		Pre-IP Medications	
		Administration of Study Medication (IP)			
		Administration of IV Fluids			
		Conscious Sedation for procedure (specify pro-	cedure):		
		Assistance with procedure(s) (please specify):			
		Other (specify)			
9.		TIGATIONAL DRUG SERVICES — (Check all that ap contact IDS@shands.ufl.edu if you check any of the boxes below for		stimate)	
		Research Protocol involves Investigational Drug	g		
		Emergency medications needed			
		Routine medications needed (includes pre-me	ds and medica	ations to treat anticipated reactions)	
	☐ Performing Conscious Sedation				
		Local anesthetics for a procedure			
10	. UF CR	C LAB SERVICES (check all that apply)			
		Lab manual for this study is available and will b	oe provided		
		Simple sample lab processing (spin, separate a	nd transfer to	aliquot)	
		Complex sample lab processing (includes use of	of additives, in	cubation or any other complex processing)	
		Urine HCG/pregnancy testing			
		Send sample for Shands Lab analysis			
		Dry ice for shipping from CRC (FedEx picks up of Dry ice can be pre-ordered 1 week in advance of use. Please ind Temporary Storage at CRC (2 weeks)			
		Plan on using CTSI biorepository (beyond 2 we	eks samnle st	orage)	

Please list all lab tests required (blood, urine, stool, spinal fluid, saliva, others), if lab kits are provided by sponsor and where analysis/final storage will take place.

Lab Tests	Kits Provided?	Analysis done at?	Same day shipping?
Example CBC	No	Shands	N/A
Example Drug Level	Yes	Central Lab	Yes

11. EC	UIP	MENT USE						
	□ Study will require the use of CRC equipment							
		Specify equipmen	t needed:					
		☐ Study Team will bring equipment to CRC						
			t and space needed:	<u>-</u>				
		(Storage of equipment w	ithin the CRC is not guaranteed, plea	se contact Nurse Manager for approv	val)			
				w to attach required docur TTACHED OR YOUR REQUE				

 ${\bf 13.\ OPTIONAL\ DOCUMENTS:\ Please\ click\ the\ button\ below\ to\ attach\ document\ files\ such\ as\ lab\ manual.\ Etc.}$

When you have completely filled out the form and attached the required documents please Click the Submit Button