

# CCTS Clinical Support Registration Form

Date: \_\_\_\_\_

Amendment?  Yes  No

**Instructions:** Fill out both sides of this form as completely as possible and submit as instructed on the back. Questions? Send email to [CCTSclinical@uab.edu](mailto:CCTSclinical@uab.edu).

IRB Title:

IRB Protocol Number:

PI Name:

PI Blazer ID:

Dept./School/Center:

Contact Name:

Phone:

Contact Email:

Coordinator Name:

Phone:

Coordinator Email:

Other Investigator Names & Blazer IDs (if UAB):

**Funding Sources** (check all that apply):

Federal agency (e.g., NIH, DOD)  
Grant Number: \_\_\_\_\_  
 Funded  Pending

Industry  
Grant Number: \_\_\_\_\_  
 Funded  Pending

Foundation  
Grant Number: \_\_\_\_\_  
 Funded  Pending

Intramural/Other  
Grant Number: \_\_\_\_\_  
 Funded  Pending

Oracle Number:

**Type of Study** (check all that apply):

Pilot  
 [Clinical Trial](#)  
 Multi-center  
 AIDS-related  
 Pediatric  
 Investigator-initiated  
 Investigational New Drug (IND)  
 Investigational Device Exemption (IDE)

**Study Details:**

Phase:  I  Ib/II  III  IV  
~ Number UAB participants: \_\_\_\_\_  
~ Duration of study in months: \_\_\_\_\_

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### Select Service(s) (check all that apply):

- [Bionutrition](#) (consultation on protocol design, development, and implementation; controlled feeding studies; monitoring participants on controlled diets; research meal services; anthropomorphic measurements; nutrient intake data; nutrition education)
- [Biorepository](#) (full spectrum of liquid biobanking from specimen intake, cataloging, storage, and retrieval; access to fresh blood and banked specimens of plasma, serum, DNA from healthy individuals; ISBER, GCP, and IRB trained staff)
- [Child Health Research Unit](#) (services tiered to meet the needs of protocols with a wide range of pediatric participants; child-friendly environment and staff with expertise in meeting physical and emotional needs of children; well-equipped exam rooms, lab services, specimen storage; assistance with study initiation, design, budget development, regulatory support)
- [Clinical Research Support Program](#) (trained and certified research coordinators [RN or non-nurse] to assist with study implementation, including adherence to regulatory requirements; communication with sponsors; protocol, data, and budget management; internal quality measures; IND/IDE applications)
- [Clinical Research Unit](#) (specially trained research nurses for inpatient or outpatient care, including assessment, testing, medication administration, IV therapy, pharmacokinetic sampling, cardiac & pulse oximetry monitoring, phlebotomy services, questionnaire administration, patient education)
- [Sample Processing \(SPAN\)](#) (specimen processing; DNA preparation; blood cell separation [PBMC preparation and cryopreservation]; preparation of lymphoblastoid lines; short-term or long-term biobanking)
- Other \_\_\_\_\_

### Please tell us more about the services you need:

**Submission Instructions:** Once you have filled out this form as completely as possible, submit via email to [CCTSclinical@uab.edu](mailto:CCTSclinical@uab.edu). Attach your protocol's schedule of events, highlighting all services you wish CCTS to provide. Also submit protocol and/or HSP and informed consent, if available. We will respond to your email within 72 hours. Questions? Submit to [CCTSclinical@uab.edu](mailto:CCTSclinical@uab.edu).