



**Lunch & Learn Special Session:
Clinical Research Town Hall Discussion**

Thursday, August 19th, 2021

Office of Clinical Research

New CDA and CTA Submission Process

All new clinical trial contracts are being processed by the Sponsored Research Services (SRS) Contract Management team at the University of Cincinnati.

Important: An executed CDA between UC and the study sponsor MUST be executed before a CTA can be negotiated by UC.

A new online submission process has been developed to support the new contracting process:

<https://redcap.research.cchmc.org/surveys/?s=CLDDCECC84>

Existing agreements executed through UC Health will continue to be managed at UC Health until their conclusion.

As always, feel free to reach out to the Office of Clinical Research for any questions

August 2021 Study of the Month

Type 1 Diabetes Study

Adults (21-40 years old) with Type 1 Diabetes (T1D)
Needed for a Research Study

What

The purpose of this research study is to determine how not eating (fasting) impacts the ability to respond to low blood sugar in people with type 1 diabetes (T1D).

Who

Adult males and females, ages 21-40, with type 1 diabetes may be eligible to participate in this research study. Participants must have had diabetes for 5+ years and must not be obese or pregnant.

Pay

Participants may receive up to \$400 for their time, effort, and travel.

Details

For more information, contact Rebecca Cason at Rebecca.cason@uc.edu or (513)-558-3427, or Jason Winnick, PhD at Jason.winnick@uc.edu or (513)-558-4437.



04-21 CCHMC IRB # 2019-0816





THERE WILL BE NO FIRST FRIDAY FOR SEPTEMBER 2021

Due to the Holiday weekend

Enjoy your Labor Day Weekend!!!

OCR Announcements:

UC Health Clinical Research Orientation and Training (CRO&T)

Thursday, September 9th, 2021
9:00 am - 3:00 pm
Virtual presentation

The last day of registration is EOB Friday,
September 3rd, 2021

Please contact Nate Harris
Nate.Harris@UCHealth.com
for information and registration

Today's Presentation:

Lunch & Learn Special Session: Clinical Research Town Hall Discussion

Brett Kissela, MD, will be on hand to moderate and address questions regarding the Office of Clinical Research move from UC Health to University of Cincinnati with the help of new, and familiar UC individuals and teams. With introductions to key contacts, roles of the departments, and where to reach out.



Brett Kissela, MD, MS

**Senior Associate Dean for Clinical Research
University of Cincinnati College of Medicine
Chief of Research Services, UC Health**



Brett Kissela, MD, MS

Albert Barnes Voorheis Professor and Chair
Department of Neurology and Rehabilitation Medicine
University of Cincinnati Gardner Neuroscience Institute
Senior Associate Dean for Clinical Research
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Office of Clinical Research



Maria Stivers
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Assisting with transition and in communication with UCH and UC teams. Primary contact for questions that cannot be directed to a specific individual or team.



Zak Johnson
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Main contact for Complion, UCH Greenphire, research in Epic and research operations. Project manager for CTMS SignalPath.



Heather Roberson
Clinical Research
Budget Administrator
vaughthr@ucmail.uc.edu
513-264-1475

Continues to work on clinical research budgets for many departments. Has access to and monitors the redcap submission form. Responsible for uploading initial budget templates and final budgets to Kuali for ALL clinical trials. Contact for any questions related to clinical trial budgets.



Macy Michael
Clinical Research Budget
Administrator
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SRS Accounting

SRS Accounting is responsible for invoicing sponsors, drawing down cash, preparing financial reports, and financial compliance.

Call SRS Accounting if you have an issue with sponsor billing, or with the ordering and distribution of ClinCards for participant payments.



John Ungruhe, Director
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Cindy Lasonczyk, Assistant Director
lasoncca@ucmail.uc.edu
513-556-4818

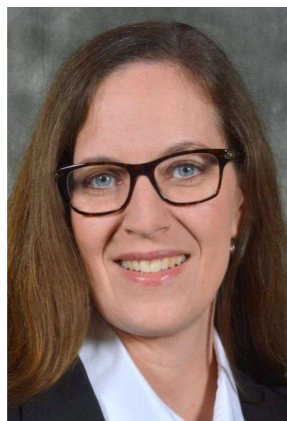
SRS Grants & Contracts



David Gearing, Director
gearidd@ucmail.uc.edu
513-556-5969

SRS Grants & Contracts Division is responsible for reviewing and negotiating the business terms of agreements, ensuring agreements are compliant with University rules, and setting up new awards in the University system

Call a member of the Industry Clinical Trials (ICT) team if you need information or help regarding contract terms or the establishment of your award account.



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Devon Robinson, Contract Specialist
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Abby Daniels
Research Unit Lead and
Sr. Asst. General Counsel
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Cynthia Dillon
Associate General Counsel
513-558-5638



Korin "Kori" Landon
Senior Legal Assistant
513-558-1726

Technology Transfer

- CDA/NDA/PIA – confidentiality agreements
- MTA – material transfer agreements
- DUA – data use agreements for de-identified data



Geoffrey Pinski
AVP, Technology Transfer
pinski@ucmail.uc.edu

CoM Accounting & Finance Pre-Award Team

- Responsible for the CoM pre-award grant submission process, from intent to submit through gathering documents and routing in Quali, responding to JIT requests, and working with SRS to set up awards.
- Also responsible for Progress Reports and NCE/Carryforward Requests
- Submissions processed through A&F Pre-Award include federal/state funding and grants that involve effort reporting, return of funds, animal/human subjects, etc.

-Generally does not include non-governmental clinical trials and investigator initiated trials.



Daniel Brummett
Executive Director

Steve Manning – Asst Director Pre-Award

- Lisa Bilinsky
- Sandra Newman
- Joanna Bordenave
- Noreen Webb-Johnson
- Julie Ison
- Stephanie Wilson



Questions?