New eDisclosure System

The Conflict of Interest Office will now be known as The Office for Responsible Outside Interests (OROI). The OROI implemented a new conflict of interest and commitment system, eDisclosure, that is designed to make submitting disclosures, research certifications and Conflict of Commitment (COC) forms easier and faster.

The UA’s new Conflict of Interest (COI) disclosure system opened July 1, 2021. The new eDisclosure page is where you can find training resources: https://research.arizona.edu/compliance/conflict-interest-program/disclosure-requirements/edisclosure-information.

Here is the link to the new system: https://edisclosure.arizona.edu/. Please see the attached email for more information.

Anyone who is listed as a COI investigator for at least one research project should have already received notifications to submit research certifications. These should be completed as soon as possible to avoid delays with IRB submissions.

notifications for annual certifications will be sent after July 12.

Please know that for their first disclosure or certification in eDisclosure, individuals will need to disclose all entities, including those previously disclosed in the legacy COI Disclosure System.

Research Certifications for which the COI review has been completed in the legacy COI Disclosure System will not need to be redisclosed in eDisclosure unless there is a modification to the project or protocol.

For questions or more information, contact the Office for Responsible Outside Interests at coi@arizona.edu or at 520-626-6406.

mPage Overview and Training

A Research Revenue Cycle mPage has been developed in Cerner to create a consistent front-end process for Banner Patient Access Services to be able to identify scheduled research encounters and ordered services that are covered by research so that the authorization, pre-registration, and registration process can be completed appropriately. Please review the attached overview. The mPage became active on February 15, 2021. Guides can be found on the Coordinator Corner website.

HIPAA Training Reminder

The UA HIPAA Privacy Program (HPP) requires all faculty, staff, and DCCs of UAHS to complete annual HIPAA certification training. The training takes about 10 minutes and provides basic information about HIPAA compliance resources at the University. Information about how to complete the training can be found on the HPP website. Additionally, the university requires all faculty, staff, and DCCs with access to university information resources to complete annual information security awareness training (ISO-500 Information Security Awareness Training Policy). The annual refresher course is approximately six (6) minutes in length, and updates employees on the latest threats, trends, and university security resources (HIPPA Annual Certification).

**Training is an essential part of a well-informed workforce. Please complete your training at your earliest convenience. If you have questions please reach out to the HIPAA Privacy Program or the Information Security Office.**

**UAHS Global HIPAA Procedures were updated in May 2021 and are available on the Research Administration website (https://research.uahs.arizona.edu/facilities-and-resources/uahs-hipaa-sops). A UA NetID is required.**

**The HIPAA Privacy Program will be offering a bi-weekly interactive seminar covering HIPAA Privacy & Security topics beginning on November 4, 2020 at 3:00 pm. Please review the list of topics (attached) or visit this link to participate in the scheduled workshops:**

https://arizona.zoom.us/j/9477665768
Research Intake Application (RIA)

New RIA for Amendments: Moving forward, please use the revised version of the amendment application attached and also available on our website. The revised form asks about the status of the project; your answers will better help our team determine how to process your application.

Informed Consent Form (ICF): To ensure we have accurate documents for coverage analysis review, we have updated the required documents for Research Intake Application (RIA) submission. Effective February 15, 2021, new and amendment study submissions to the RIA require that the submitted ICF template includes tracked changes with any required Banner or UA language.

Protocol Amendments: It is very important that protocol amendments be submitted through the RIA process as soon as you receive them. Protocol amendments undergo a review and update of the coverage analysis (CA) and the OnCore calendar/financials. Both can be completed concurrently with IRB review and approval. IRB approval is not required for RIA submissions. This will allow us to update your OnCore calendar so it is ready for release as soon as IRB approval is received.

Clinical Trials Website: Please be sure to “opt-in” to having your study published on this website. This can be found on pages 5-6 of the Research Intake Application (RIA). It is a great way to build collaboration within the research community and for potential study subjects to find studies. We are adding a “COVID-19 Research” heading for all studies associated with this category. If your study is not currently listed, please contact our office at crc@email.arizona.edu.

RII Research Restart Checklist for COVID Research

NOTE: Banner updated their Research Guidance (attached) for research studies on 05/1/2021. Access to their facilities may impact the approval of research studies.

The UA has transitioned to Phase 5 of the Research Re-Start Plan, which means that restart checklists are no longer required. Information on Phase 5 is located here.

- For COVID studies occurring in Banner space, approval from the UA-COVID committee is required. If you have approval, please indicate this in the abstract section. If you are not sure you have approval, please email Anna Valencia (Phoenix).

Outlook Contacts to Add to Avoid Missed Messages

Research Administration serves investigator teams across UAHS through a wide range of pre- and post-award activities. Frequent and timely contact keeps those processes moving, and that means a large volume of email. Adding Research Administration email addresses to your Outlook contacts can ensure that time-sensitive messages don’t end up unnoticed in your Junk Email folder.

Updating contacts is especially important for research teams who are working with OnCore, since the system frequently sends automated messages in batches from OnCoreSupport@email.arizona.edu. Some users have reported that messages about OnCore have ended up in their spam folder. CRC@email.arizona.edu, UAHSContracts@email.arizona.edu, and ResearchApp@email.arizona.edu are additional email addresses that should be added in Outlook due to the high volume of messages they send.

Contacts can be added from a new message by right-clicking on an email address and selecting Add to Outlook Contacts, or by going to the Contacts (or People) tab in your left sidebar and selecting New Contact from the top ribbon.

Research Administration maintains a Departmental Contacts page that includes a comprehensive list of email addresses for all of the work groups that could be in communication with you. Sponsors and clinical partners are also contacts to consider adding in Outlook.
OnCore, Training and Individual Consultations

Starting in July, OnCore Support will replace its regular office hours with self-service scheduling for support sessions through Microsoft Bookings. Individual consultations are available in a HIPAA-compliant Zoom environment in case specific examples from your OnCore study are discussed. A Zoom link will be provided in your email confirmation once you schedule your session.

Monthly trainings for new OnCore users will continue on the same schedule, with training sessions held the first full week of each month (occasionally adjusted for holidays or other events). Available trainings are posted one to two months in advance.

The OnCore website provides information about scheduled trainings and individual support sessions on the Training and Consultations page. Please feel free to sign up if you would like first-time training, a refresher training, or one-on-one OnCore help!

To register for the next training sessions, please complete an OnCore Confidentiality Agreement and send your training request to OnCoreSupport@arizona.edu. The next trainings are scheduled as follows:

- Introduction to OnCore and Calendar Validations
  Tuesday, July 13, 1 pm – 3 pm, August 3, 1 pm – 3 pm
- Subject Management Training
  Wednesday, July 14, 1 pm - 3 pm, August 4, 1 pm – 3 pm
- Regulatory Training
  Thursday, July 15, 10 am – 12 pm, August 5, 10 am – 12 pm

If you have changed departments or need to have an additional role added to your OnCore Profile (regulatory, study coordinator, etc.), you will need to submit an updated OnCore Confidentiality Agreement to OnCoreSupport@arizona.edu prior to the role being added. Additional training may be required.

We are also available to attend department or research unit meetings. This is a great way to receive direct support for your team’s research studies and ask specific questions for OnCore Support. Please email us at OnCoreSupport@arizona.edu to schedule a session.

Subject entry can begin when calendars have been validated, IRB documents have been uploaded, and the study has been opened to accrual by the regulatory team. Studies will need a fully executed or signed contract prior to being opened to accrual in OnCore (as applicable). Please be sure to enter each subject’s country and zip code on the Subject Demographics page.

All subject visits MUST be checked in/logged into OnCore within 24 hours of study visit.

Year-One Feedback Sought for OnCore Clinical Trial Management System (CTMS) and OnCore Support

OnCore CTMS launched as a UAHS enterprise system in April 2020. Due to the scope of the project, the launch consisted of a rolling implementation plan, involving close work with each department to bring their protocols into the OnCore environment. That means many OnCore users have been using the system for just over a year.

Prior to the enterprise launch, OnCore had been in use at the UA Cancer Center since 2007, but the launch to a wider segment of the campus meant changes for UACC users as well.

The project's success was also due to each and every clinical trial investigator, manager, and coordinator who took the time to learn about OnCore and its associated workflows. For that reason, OnCore Support would like to invite those users to provide their feedback on what’s worked and provide feedback and suggestions for improvement.

In the coming days, a survey will be shared via the OnCore listserv to invite your anonymous feedback on the system and its support team. The data we collect will provide feedback for us to improve our services and steer the future of OnCore Support. Your time, consideration, and candor are appreciated during the survey period. The survey will be open through Friday, July 23, and responses from all current or recent OnCore users are welcome. If you have any questions about the survey, please feel free to reach out to OnCoreSupport@arizona.edu.
OnCore, Training and Individual Consultations, continued

UAHS Sign-off in OnCore: This sign-off is done by Research Administration upon completion of the coverage analysis (CA), budget, and fully executed contract (if applicable) and receipt of the IRB approved ICF(s).

Regulatory in OnCore (REQUIRED FOR ALL STUDIES):

- **New Studies**: Please upload your approved IRB documents (approval letter, protocol, and approved ICF(s)) into OnCore. Documents should be uploaded using the PC Console (PC Console > Reviews > IRB). Please verify that the NCT Number has been added (PC Console > Main > Details).

- **Amendments**: Protocol amendments, IRB approval letters, and the newly approved ICF(s) (as applicable) need to be uploaded into OnCore using the PC Console. The amendment IRB approval date needs to be entered. IRB approval of the protocol amendment will help the OnCore Support team know when to release the updated calendar for the protocol amendment (as applicable).

- **Personnel Changes**: Please be sure to update any personnel changes in OnCore, update IRB approval/closure dates and upload IRB approval documents (approval letters, ICFs, etc.)

- **Study Closure**: Upload IRB closure notice, change the study status to “IRB Study Closure”, and enter the study closure date.

Cerner & OnCore: OnCore is now able to push “On Study” subject information to Cerner. This will add a notification on the blue banner to the patient’s medical record that they are enrolled in a UA clinical trial. All active protocols with active subjects for have been pushed over to Cerner. Once a subject is marked as “On Study” be sure to check Cerner to verify that the blue banner appears. If it does not appear, verify that the first and last name, date of birth, and MRN all match. Phase II of the OnCore/Cerner interface is for Cerner demographics information (MRN, Name, DOB, gender, race, ethnicity, and address) to push to OnCore. This is currently under development.

Next Steps

- During the next several months:
  - We are continuing to work on entering study budgets into OnCore. This will aid with invoicing sponsors and tracking study payments. We will reach out to departments when we are ready to schedule training.
  - Provide overview and training on running reports from OnCore
- Implementation of the eRegulatory Management System was completed in March. A formal launch date is still being determined.

Please email us at OnCoreSupport@arizona.edu with questions, or for additional help.
Banner Hospital Billing Update

Banner Hospital billing for the months of October 2017 – May 2021 have been reviewed and sent out to the corresponding UA Departments via UABox Health.

- Please process payment promptly. Payments are due 30 days from receipt of the billing. If there are any discrepancies, please email ctfinance@arizona.edu for assistance.

- When submitting backup to FSO, please only redact the patient name and date of birth if applicable. All other information should be left visible. Please see example below (this is a fictional bill with no HIPAA information)

- Please send an email to ctfinance@arizona.edu with your DV payment information.

- Please do not Closeout and FPC any account balances if your clinical trial protocol reflects Banner services. If you are unsure, please work with your Study Team for confirmation.

- Payments need to be processed within 30 days of billing receipt.

Please use GL Code #4215 for all payments and purchase orders to Banner Health. This GL code was created to capture all research related expenses for ‘Various clinical trial procedures, i.e. imaging, venipuncture, labs, exams, etc.’. This allows for smoother account reconciliation and reporting.

Billing Compliance Process for Clinical Trials Purchasing BH Services

The University of Arizona is obligated to log all study visits into OnCore. Study visits must be logged within 24 hours of occurrence whenever Banner Health (BH) services are utilized for a research study (i.e. medical imaging, ECG, clinic visits, etc.).

These services are typically scheduled via Cerner on behalf of the research patient. ALL study visits that include BH services MUST be logged into OnCore within 24 hours.

- This includes research-related AND routine/standard of care.

- UA Coverage Analysis (CA) provides detailed information for billing designations. Study calendars in OnCore reflect these billing designations. A copy of the CA is uploaded into OnCore for the study team’s reference.

- This process helps to ensure that bills are routed to the correct payer and helps to protect a study subject and alleviate incorrect billing.

BHRF reviews and validates all charges logged into OnCore against what has been billed in Cerner. Charges are then generated and billed to the research study or subject’s insurance as verified by the coverage analysis.

If you have questions regarding the OnCore calendar, contact OnCoreSupport@arizona.edu.

Questions regarding the coverage analysis? Contact Research Administration at crc@arizona.edu.
COVID-19 Research and Sample Request Guide

The University of Arizona research community has been actively studying patients infected with COVID-19 in hopes of learning more about the virus, its pathogenesis and possible treatments.

As part of these efforts, the University of Arizona Health Sciences Biorepository created the COVID-19 Research and Sample Request Guide (attached) for researchers using biospecimens in COVID-19 studies.

Investigators wishing to initiate a COVID-19 study that would require biospecimen collection should contact Dr. Sairam Parthasarathy at sparthta1@arizona.edu for patient access.

The Health Sciences Biorepository provides an electronic universal consent, along with a REDCap database and linkage to electronic medical data stored in Cerner for each subject.

Please submit any request for COVID-19 samples at https://biobank.uahs.arizona.edu or http://redcap.link/covid19request.

To review available samples in the biobank, please see the Biorepository Summary.

For more information, please review the attached guide or contact Dr. David Harris, Director of the Health Sciences Biorepository at davidh@arizona.edu.

Sonora Quest Laboratories Account Set-up and/or Care360 User Request

Email requests to: ctfinance@arizona.edu

Please include the following information with your request:
Name, Job Title, Net ID, UA Email, Phone and Fax numbers, Physical Work Address, Department, SQL Account Number (if known)

Study Close-out with IRB and Final Study Payments

Once your study has been closed with the IRB, remember to enter the closure date into OnCore. Please be sure to work with your business office to verify all payments to vendors have been issued and that all invoiceable items have been sent to your study sponsor (as applicable). Clinical trial contracts have a specific window to complete these tasks. If you are unsure or have questions, please contact our office at ctfinance@arizona.edu or crc@.arizona.edu.

UAHS Clinical Research Professionals (CRP) Group Meeting

If you are new to the University of Arizona Health Sciences (UAHS) research community and/or would like to keep up with the ever-evolving changes in UAHS research, please feel free to attend the monthly CRP group meetings. Meeting time and location changes from month to month and an email reminder is sent out prior to the monthly meeting.

To add your name to the listserv, please send an email to clinicalresearchcoordinators-request@lists.arizona.edu with "SUBSCRIBE" in the subject line.

"Each department/division is responsible for sending at minimum one delegate to attend the CRP meeting. If a department/division cannot attend, then the manager/supervisor will need to attend a makeup session to review topics covered in the CRP meeting."**

We welcome your feedback!! Please let us know if there are specific topics that you would like to have covered at upcoming meetings. Please send an email to vphs-cro@arizona.edu.

CRP meetings will now be held every other month starting with the May meeting.

The next scheduled meeting is Thursday, July 22, 2021, from 3:00 pm - 4:30 pm via Zoom.

Join Zoom Meeting: https://arizona.zoom.us/j/81488925948
Meeting ID: 814 8892 5948
One tap mobile
US: +16027530140,81488925948#

CRP Group upcoming meeting schedule:

<table>
<thead>
<tr>
<th>Date</th>
<th>Time</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thursday, Jul 22, 2021</td>
<td>3:00pm - 4:30pm</td>
</tr>
<tr>
<td>Wednesday, Aug 18, 2021</td>
<td>12:00pm-1:30pm</td>
</tr>
<tr>
<td>*Review of new eIRB system</td>
<td></td>
</tr>
<tr>
<td>Wednesday, Sep 15, 2021</td>
<td>12:00pm - 1:30pm</td>
</tr>
<tr>
<td>Thursday, Nov 18, 2021</td>
<td>3:00pm - 4:30pm</td>
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</tbody>
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GENERAL INFORMATION AND RESOURCES

UAHS Research Administration provides guidance and assistance with the following:

- Our website: https://research.uahs.arizona.edu/
- Coverage Analysis (CA) and Clinical Trial Budget development/ negotiations: contact: crc@arizona.edu
- Contracts (CDAs, NDAs, CTAs, amendments, data use, incoming MTAs): contact: UAHSContacts@arizona.edu
- Regulatory contact: regulatory@arizona.edu or schedule 1:1 session
- Post-Award Clinical Trial Accounting and Auditing: contact: CTFinance@arizona.edu

UAHS Project Status Report: https://research.uahs.arizona.edu/facilities-and-resources (UA NetID Login required)

Research Intake Application (RIA):
Applications and required documentation should be emailed to ResearchApp@arizona.edu. Instructions and application forms can be found here:
http://research.uahs.arizona.edu/clinical-trials/research-intake-form

If you have questions, email Research Administration at crc@arizona.edu.

UAHS OnCore Support: OnCoreSupport@arizona.edu or https://research.uahs.arizona.edu/oncore or schedule 1:1 session (calendar validations, subject management, regulatory, IT support)

ClinicalTrials.gov Assistance:
- Non-cancer studies: Kirsten Anderson, regulatory@arizona.edu or (520) 621-6417
- Cancer studies: Amy Selegue, UACC-NCTN@uacc.arizona.edu, (520) 626-0301

UA HIPAA Privacy Office: Contact PrivacyOffice@arizona.edu or (520) 621-1465

UAHS Global HIPAA Procedures: https://research.uahs.arizona.edu/facilities-and-resources/UAHS-hipaa-sop’s (NetID Login required)

IRB Training Opportunities
The IRB offers training on a variety of topics each month. This is a great way to stay updated on current processes and have your questions answered. The list of upcoming sessions is located on the IRB website with instructions for signing up through UAccess EDGE Learning.
https://rgw.arizona.edu/compliance/human-subjects-protection-program/irb-training-opportunities

REDCap Questions/Training: Contact redcap@arizona.edu

Data Warehouse Information: https://research.uahs.arizona.edu/clinical-trials/resources#data

UA Clinical and Translational Science (CATS) Research Center: http://cats.med.arizona.edu


Banner Badge Request: Contact clinicalresearch@arizona.edu

Banner Cerner Help: Contact the Banner IT service desk at (602) 747-4444 or in Tucson, call (520)-694-HELP (4357). Select Option 6 for assistance with Multi-factor Authentication.

Cerner Access/Training: Contact your department’s assigned Banner Health Clinical Trial Senior Manager.
https://research.uahs.arizona.edu/clinical-trials/cerner

Sonora Quest Laboratories Account Set-up: email request to ctfinance@arizona.edu


SQL Care360 Training: Contact the Customer System Team at (602) 685-5465 or SQLCustomerSysem@SonoraQuest.com to schedule training. Please be sure to include your SQL departmental account number when requesting training.