

## TRAcS – Menu

Below is a list of the options available in the UMCCTS TRAcS along with the files to be uploaded and required information. Files without notations or information listed without notations are required. Notations: (opt) = optional; (if applic) = if applicable

“Pass through” means that UMCCTS staff will forward the request to the appropriate service provider.

Some selected options will direct the user to the affiliated web pages to make their request.

**COVID FastTracs** – has been disabled at this time. Enter any COVID related request in the usual manner, but indicate that the request is COVID-related (and RADx related, if applicable).

TRAcS option	Documents to Upload / Required information	Service Provider Responsible
<b>Protocol Start Up or Implementation</b>		
<b>Biorepository &amp; Tissue bank</b> <ul style="list-style-type: none"> <li>• Obtain samples from the Biorepository/Tissue Bank</li> <li>• Provide samples to the Biorepository/Tissue Bank</li> <li>• Letter of support</li> <li>• Consultation</li> <li>• Request a quote</li> <li>• Other</li> </ul>	Draft Letter of Support (if applic, opt)	Biorepository & Tissue bank staff  Email notification to core by TRAcS Administration
<b>Clinical Research Center</b> <ul style="list-style-type: none"> <li>• Clinical (blood draws, blood pressure etc.)</li> <li>• Space (clinic rooms, freezer storage, etc)</li> <li>• Data management (entry, etc)</li> <li>• Full study coordination</li> <li>• Update to an existing service agreement</li> <li>• Pricing only - estimates for a proposed project</li> <li>• Other</li> <li>• Not sure, want to ask a question</li> </ul>	Protocol or study plan Informed consent form Investigator Brochure (opt) Lab manual (opt)	CRC staff
<b>Office of Clinical Research (OCR) : Contracts &amp; Budgeting</b>	Space for additional document (opt)	
<ul style="list-style-type: none"> <li>• Have/want a new Data Use Agreement (DUA)</li> </ul>	Email requesting contract agreement IRB approval letter Draft DUA (if applic, required)	OCR - Research Compliance Contracts
<ul style="list-style-type: none"> <li>• Have/want a new Confidentiality Disclosure Agreement (CDA) or Non-Disclosure Agreement (NDA)</li> </ul>	Confidentiality Disclosure Agreement (CDA) (if applic) Email from sponsor (opt)	OCR - Research Compliance Contracts

TRAcS option	Documents to Upload / Required information	Service Provider Responsible
<ul style="list-style-type: none"> <li>Have/want a new Clinical Trial Agreement (CTA)</li> </ul>	Protocol or study plan Informed consent form Sponsor budget NCT number IDS Pharmacy Budget Draft Clinical Trial Agreement CRC service agreement (if applic) Proposal Routing Form Investigator Brochure (opt) Lab manual (opt) Email from sponsor (if applic) IND or IDE Waiver (if applic)	OCR – Preaward OCR - Research Compliance Contracts
<ul style="list-style-type: none"> <li>Have a newly awarded clinical research grant</li> </ul>	Protocol or study plan Informed consent form NCT number Notice of Grant Award Budget template with schedule of events or schedule of events PIN Report (account number) IND or IDE Waiver (if applic) Investigator Brochure (opt) Lab manual (opt)	OCR - Preaward
<ul style="list-style-type: none"> <li>Have/want a new federally-funded cooperative group trial</li> </ul>	Protocol or study plan Informed consent form Master/ Clinical Trial Agreement(s) Study fund sheet NCT number National coverage analysis (opt) IND or IDE Waiver (if applic) CRC service agreement (if applic) Proposal Routing Form or PIN Report (account number)	OCR - Preaward
<ul style="list-style-type: none"> <li>Have a new internally-funded clinical research trial</li> </ul>	Protocol or study plan Informed consent form Budget template/schedule of events NCT number Account number for invoices IND or IDE Waiver (if applic) CRC service agreement (if applic)	OCR - Preaward
<ul style="list-style-type: none"> <li>Have a protocol or contract amendment</li> </ul>	Protocol/calendar - tracked Budget/contract – tracked	OCR - Preaward
<ul style="list-style-type: none"> <li>Need to revise/correct OnCore information</li> </ul>	Protocol/calendar - tracked Budget/contract – tracked	OCR - Preaward
<ul style="list-style-type: none"> <li>Would like to obtain a Certificate of Confidentiality</li> </ul>	IRB Approval letter Informed consent form UMMS CoC letter signed by the PI	OCR - Preaward

TRAcS option	Documents to Upload / Required information	Service Provider Responsible
<ul style="list-style-type: none"> <li>Want to submit a clinical research account closure request</li> </ul>	Account closure form (opt) Reconciliation/tracking in any format Sponsor payment history (opt) PeopleSoft Reports/Summit Screen	OCR - Preaward
<ul style="list-style-type: none"> <li>Need review/processing an Institutional Prior Approval (IPA) for a change in budget or extend study period</li> </ul>	IPA form	OCR - Preaward
<ul style="list-style-type: none"> <li>Need help with the Conquering Diseases clinical study opportunities portal</li> </ul>		TRAcS Navigator
<ul style="list-style-type: none"> <li>Need new user access for Epic</li> </ul>		OCR - Postaward
<ul style="list-style-type: none"> <li>Don't know/ I want to ask a question</li> </ul>	Space for additional document (opt)	TRAcS Navigator
<b>Investigational Drug Services</b> <ul style="list-style-type: none"> <li>Pricing for clinical trial support services</li> <li>Set up to dispense drugs for a clinical study</li> <li>Assistance with obtaining or renewing a Massachusetts Controlled Substances Registration (MCSR) for clinical research</li> </ul>	Protocol or study plan Informed consent form Pharmacy manual (opt) Investigator Brochure (opt)	IDS Staff
<b>IRB consult or question</b> <ul style="list-style-type: none"> <li>Extramural Institutional Certification</li> <li>IRB question / consult</li> </ul>		TRAcS Administration
<b>Clinical Trials.gov assistance</b> <ul style="list-style-type: none"> <li>I need an account for myself</li> <li>I need an account for someone else</li> <li>I need help with entry / maintenance / navigation</li> <li>Assistance determining whether an NCT number is required for my study</li> <li>Other</li> </ul>		OCR - Research Compliance Contracts
<b>Protocol Review Committee</b>	Protocol or study plan Informed consent form Feasibility checklist	TRAcS Navigator
<b>Recruitment Resources</b>		
<b>Connect to Conquering Diseases or ResearchMatch research volunteer registry members about participation in a study</b> <ul style="list-style-type: none"> <li>Access to volunteer registry for recruitment</li> </ul>		TRAcS Navigator
<b>Community Engagement</b>		Directed to Community Engagement forms  Email confirmation by TRAcS Administration

TRAcS option	Documents to Upload / Required information	Service Provider Responsible
<b>Informatics (Recruitment Core, Data ScienceCore, TriNetX)</b> <ul style="list-style-type: none"> <li>• Recruitment Core - Access/develop tools and alerts for recruitment and cohort identification</li> <li>• Data Science Core - Obtain data for research from the Data Lake</li> <li>• Data Science Core - Request a research informatics consultation</li> <li>• Request a TriNetX account</li> </ul>		Directed to appropriate IT forms  Email confirmation by TRAcS Administration
<b>Interpreter or Translation Services for Research</b> <ul style="list-style-type: none"> <li>• Over the phone interpreter services</li> <li>• Video interpreter services</li> <li>• In-person interpreter services</li> <li>• Translation services</li> </ul>		TRAcS Navigator
<b>Research Navigator Service (consult)</b> <ul style="list-style-type: none"> <li>• Consult with the Navigator</li> <li>• Access to volunteer registry for recruitment</li> <li>• Information about the Trial Innovation Network (TIN) infrastructure and resources for multi-site trials</li> <li>• I'm not sure, I want to ask a question</li> </ul>		TRAcS Navigator
<b>Other Services or Requests</b>		
<b>Bioinformatics</b>		Email notification to Bioinformatics by TRAcS Administration
<b>CCTS Membership</b>		TRAcS Administration
<b>Education / Training / CRPG</b> <ul style="list-style-type: none"> <li>• I would like to take a Study Coordinator course</li> <li>• I would like to request individual Education or Training for me or my team</li> <li>• Include me on the Clinical Research Professional Group (CRPG) email list</li> </ul>		CCTS Educator
<b>Find Funding Opportunities</b> <ul style="list-style-type: none"> <li>• Interested in learning about UMCCTS funding opportunities</li> <li>• Interested in collaborating on UMCCTS pilot grant</li> <li>• Interested in other UMass funding opportunities</li> <li>• Interested in external funding opportunities</li> <li>• Other</li> </ul>		CCTS Administration
<b>Library Services</b>		Directed to library forms

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<b>Massachusetts Medical Device Development Center (M2D2)</b> <ul style="list-style-type: none"> <li>• I am interested in M2D2 funding opportunities</li> <li>• I am interested in M2D2 educational events</li> <li>• I am interested in talking about renting M2D2 space</li> <li>• I am interested in learning about M2D2 resources and services</li> <li>• Other</li> </ul>		CCTS Administration
<b>Quantitative Methods Core</b> <ul style="list-style-type: none"> <li>• clinical research support in biostatistics, experimental design, and data management</li> </ul>		Directed to QMC forms  Email confirmation by TRAcS Administration
<b>Small Molecule Screening Facility</b> <ul style="list-style-type: none"> <li>• SMSF Equipment use</li> <li>• Design of screening assays</li> <li>• Letter of support</li> <li>• Consultation</li> <li>• Request a quote</li> <li>• Other</li> </ul>	Draft letter of support (opt)	Ann Han or Margaret McManus – pass through to SMSF
<b>Science Participation Research Center (SPRC)</b> <ul style="list-style-type: none"> <li>• Increase engagement of special populations in translational research through tailored, culturally responsive strategies</li> </ul>		SPRC staff  Email notification to SPRC by TRAcS Administration
<b>Study Audit (Internal, QA)</b>		CCTS Educator
<b>Umbilical Cord Blood Core</b>		Cord Blood Core staff  Email notification to core by TRAcS Administration
<b>Miscellaneous Study Conduct (e.g. lab certs, MCSR, and sponsor to eIRB etc)</b> <ul style="list-style-type: none"> <li>• I would like to a copy of the lab certifications</li> <li>• I would like information about MCSR (Massachusetts Controlled Substances Registration)</li> <li>• Add sponsor to eIRB and /or OnCore</li> <li>• Other</li> </ul>		TRAcS Navigator
<b>I can't find what I need</b>		TRAcS Navigator

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**For more information:**

Any questions can be directed to Clinical Research Navigator [ann.han@umassmed.edu](mailto:ann.han@umassmed.edu) who will be happy to assist you.

Questions related to OCR can be directed to: [clinicalresearch@umassmed.edu](mailto:clinicalresearch@umassmed.edu)