**Clinical research resources**

INVESTIGATOR-INITIATED PIPELINE

1. Assessment
	1. [Draft protocol and study related documents](https://research.vcu.edu/forms/) (located under HRPP/IRB tab)
2. Start-up
	1. Study activation
		1. [Budget/cost coverage analysis](https://research.vcu.edu/human-research/clinical-resources/)
		2. [Subject injury language](https://research.vcu.edu/media/office-of-research-and-innovation/secure/proposalsandawards/rams-spot/osp_subject_injury_language_review_process_for_industry.pdf)
		3. [PROC/VCUHS intake form](https://onetrac.vcu.edu) or [PRMC](https://www.masseycancercenter.org/research/~/link.aspx?_id=ee49e95faa8b44d09b6e89d8e3b48b57&_z=z)
		4. [Ancillary review](https://research.vcu.edu/integrity-and-compliance/compliance/regulatory-committees/)
		5. [IRB review](https://research.vcu.edu/human-research/hrppirb/activities-requiring-irb-review/)
		6. [Clinical trial registration](https://cctr.vcu.edu/support/consultation/clinical-trials-gov/)
	2. Study set-up
		1. Regulatory files ([SiteVault](https://research.vcu.edu/human-research/regulatory-affairs/veeva/))
		2. [Staff training/delegation](https://research.vcu.edu/training/citi-training/)
		3. [OnCore](https://cctr.vcu.edu/support/informatics/oncore/)
		4. Site initiation
		5. [Regulatory filing (IND/IDE)](https://research.vcu.edu/human-research/regulatory-affairs/)
3. Conduct
	1. [Recruitment](https://cctr.vcu.edu/support/study-participation-and-recruitment/)
		1. Advertising
		2. Recruitment
		3. Screening
		4. Consenting
	2. Management
		1. [OnCore](https://cctr.vcu.edu/support/informatics/oncore/)
		2. [Ordering and scheduling](https://research.vcu.edu/human-research/ids/)
		3. [Biospecimen management](https://pathology.vcu.edu/research/clinical-pathology-cprs/)
		4. Participant follow-up
		5. [Reimbursement/compensation](https://procurement.vcu.edu/i-want-to/pay-an-individual/compensate-a-research-participant/)
		6. [Research billing](https://intranet.vcuhealth.org/sites/clinical-research/SitePageModern/80467/study-billing)
		7. Participant records
		8. [Data collection](https://cctr.vcu.edu/support/research-navigation/starting-data-collection/)
		9. Safety monitoring
		10. Re-consenting
	3. Oversight/maintenance
		1. [Site file maintenance](https://research.vcu.edu/human-research/regulatory-affairs/veeva/)
		2. [IRB continued review](https://research.vcu.edu/human-research/hrppirb/other-submissions-and-monitoring/)
		3. [Amendments](https://research.vcu.edu/human-research/hrppirb/other-submissions-and-monitoring/)
		4. [Reportable events](https://research.vcu.edu/human-research/hrppirb/other-submissions-and-monitoring/)
		5. [Regulatory reporting](https://research.vcu.edu/human-research/regulatory-affairs/)
		6. [Data management](https://ts.vcu.edu/about-us/information-security/data-management-system/)
		7. Data query resolution
		8. Study monitoring/audits
4. Closure
	1. Close out visit
	2. [IRB closure](https://research.vcu.edu/human-research/hrppirb/other-submissions-and-monitoring/)
	3. Archiving records
	4. [Clinicaltrials.gov](https://cctr.vcu.edu/support/consultation/clinical-trials-gov/) results reporting
	5. [Regulatory Closeout](https://research.vcu.edu/media/office-of-research-and-innovation/clinical/Regulatorychecklistforstudycloseout.docx)

For assistance, please contact your appropriate school for assistance:

* Massey Cancer Center: masseyactpm@vcu.edu
* School of Medicine: SOMCT@vcuhealth.org