



An NCI-designated Comprehensive Cancer Center

Strategies for Expansion and Efficiency of Biospecimen Management for Multi-site Investigator Initiated Trials (IITs)

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Background

Over the course of 10 years, the Research Biofluid Management Unit (RBMU) of the Clinical Trial Office (CTO) at NYU Langone Health (NYULH) Perlmutter Cancer Center (PCC) has supported the exponential growth of multi-site investigator-initiated trials (IITs) from 3 to 17 research studies, with patient enrollment growing from 85 participants in 2016 to 580 participants across trials in 2024. To support such growth, RBMU biospecimen management needed to be streamlined and standardized across sites. Creation of sustainable and efficient biospecimen management workflows is crucial to reduce the loss of valuable human specimens and maintain reliable pre-analytical conditions for downstream biospecimen analysis at NYU Langone Health research laboratories.

Goals

- Identify cost-effective and efficient solutions that would enable expansion of IIT activity across multiple sites at a national scale.
- Implement operational workflows that reduce time of supply provision to sites.
- Provide thorough processing/shipping instructions in a standardized manner to reduce deviations and errors upon sample submission to NYU Langone laboratories.
- Adapt for growing scope of processing techniques and requirements commonly found in IIT research.

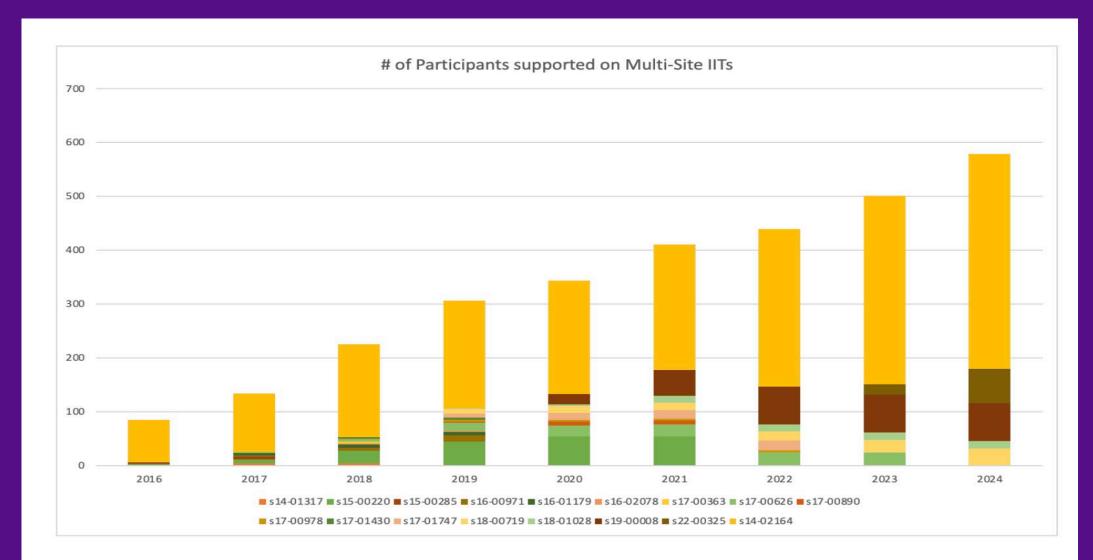


Figure 1. Patient accrual growth on multi-site IITs from 2016 to 2024

Solutions and Methods						
 Consultation with PIs during protocol 						•
development for Specimen Management						
 Protocol-Specific Lab manual creation 						
 Pre-labeled kit provision to external sites 						•
 Redcap order form request system 						
 Redcap specimen submission system 						
 Increase specialty FTEs for overnight specimen 						•
shipments and PBMC isolation						
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Outcomes

- Increased efficiency of specimen intake process with pre-labeled specimens and electronic manifests
- Decreased time of kit order fulfillment from 3 weeks to 1 week (on average, +/- 7 days)
- Minimized errors in specimen collection and
- shipment with use of lab manual, provided kit
- supplies, pre-printed shipping labels, and shipping supplies
- Enabled growth of IIT support, increased range of specimen processing techniques, absorbed PBMC isolation processing previously
- outsourced.

Lessons Learned & Future Directions

plementation of standard instruction, supplies, and becimen management for IITs is crucial for reliable research atcomes and downstream biospecimen analysis. By inimizing the risk of deviations due to specimen ocessing/shipping errors at external sites, we were able to reamline support for IITs at our cancer center and enable verall expansion of multi-site activity. Going forward for ontinued expansion, we may look to outsource kit provision transfer the burden of creating "homemade" specimen ollection and processing kits from our staff to a third party. It II also be necessary to integrate external specimen abmission with current RBMU LabVantage LIMS specimen anagement programs currently in the final stages of evelopment and implementation this year.