

National Institutes of Health (NIH) Biomarkers for Vascular Contributions to Cognitive Impairment and Dementia Consortium (MarkVCID)

Refer to section 3.4.4 in the Manual of Operating Procedures before initiating the process.

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Communication and Approvals							
Enrolling site							
☐ Confirm participant's willingness to transfer to the receiving site							
☐ Contact the receiving site PI to discuss participant transfer and timeline.							
 Receiving site PI confirms site has resources for remaining study visits Clinical and administrative personnel MRI availability, biospecimen storage facilities Other supplies 							
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Both enrolling and receiving sites							
 Follow site's IRB and regulatory requirements to transfer or receive the participant (Check with your institution for specific regulations.) 							
☐ Notify the CC of IRB approvals							
Enrolling site							
☐ Notify participant of 1) approved transfer 2) new study team's point of contact							
 Share participant contact information with the receiving site and work with them to ensure the participant is fully informed and the transition is seamless (Use encrypted email or other HIPAA-compliant communication channel for sharing information.) 							
Receiving site							
☐ Contact and consent the participant							
 Notify the enrolling site and CC of the reconsent status via email and record in the Participant Transfer Form in the EDC 							
Participant Data Management							
Enrolling site							
☐ Close out outstanding data entry and queries							
☐ Upload imaging files							
☐ Record all biosamples associated with the participant							
$\ \square$ Transfer participant source documents to the receiving site through secure file system							
Scan biosamples out of the virtual biorepository and ship participant biosamples to the receiving site (Visit MarkVCID2 Protocols & Resources > Biospecimen Collection Best Practices & Shipping Procedures dropdown menu > "Shipping Human Biospecimens Guideline" and "Biorepository Sample Tracking Instructions Manual")							
☐ Initiate Participant Transfer Form in the EDC							
☐ Enrolling site ID, reason for request, last completed visit, next visit window date							
Receiving site							

Comp	olete Participant Transfer Form in EDC
	Receiving site ID; confirm transfer of the participant ID in EDC, reconsent prior to transfer, and that research data records were received

Receive ship	ment and scar	n biosamples int	to the virtual	biorepository

Schedule participant's next study visit