

Managed by **Duke** Clinical & Translational Science Institute

The MURDOCK Study Community Registry and Biorepository is a 12,526-participant community-based longitudinal cohort recruited from a 20-Zip Code region in the Southeastern United States (U.S.) that is centered in the city of Kannapolis, NC and encompasses Cabarrus County, NC.

Creation of the cohort was funded by a gift to Duke University from the David H. Murdock Institute for Business and Culture, with operational support from Duke's Clinical and Translational Science Award (CTSA) grant (UL1TR002553) and the Duke Clinical and Translational Science Institute (CTSI).

Consenting participants complete a baseline health questionnaire at enrollment, as well as a brief physical exam and collection of blood and urine. Consent includes permission to access to information from medical records, storage of collected samples in the biorepository, access to collected data and biospecimens for future approved research studies and contact regarding new research study opportunities.

Data have been organized into "storefronts" that summarize characteristics of a population of research interest as well as available data and samples for that population. The following sections summarize the sources of data in the MURDOCK Study database, as well as important descriptions and definitions to help understand the data presented in the "storefronts".

1 Participant self-reported data at baseline. The baseline questionnaire collects contact information, current residential street address, and primary physician; alternate contact information; date and place of birth; demographics; current or past diagnosis of 34 medical conditions; menopausal status in women; medications, vitamins and supplements; dietary and physical activity assessment; hours of sleep per night; tobacco and alcohol use; second-hand smoke exposure; and selected PROMIS® participant-reported outcomes domains. Socioeconomic data collected at baseline included marital status, highest level of education of participant and participant's mother and father, employment status, mother's and father's occupations, housing (type, how paid for, number of adults and children in the household) and total household income. In addition, a brief physical exam (vital signs, height, weight, and waist circumference) was conducted at enrollment.

Medical conditions: "Do you have, or have you ever had, any of the following [medical conditions]?" (yes, no, don't know). Counts are unique participants reporting yes to specific condition. Medications: "Please list any pharmaceutical and/or natural medications (including vitamins) that you are currently taking." Data are captured in free-text format as written by the participant and coded using RxNorm. Summary metrics are based on everything reported. Top 5 reported medications are limited to reported prescriptions.

2 Biorepository samples. Blood was collected at baseline and processed into the following specific samples: whole blood in EDTA for DNA extraction, whole blood in PAXgene for RNA extraction, plasma, serum and buffy coat in cryovials. Urine was collected and aliquoted in cryovials. Sample collection was not done systematically for MURDOCK enrollees; however, some nested sub cohorts and other studies enrolling MURDOCK registry participants include sample collection at follow up time points. All samples are stored at -80°C in a central biorepository current managed by Fisher BioServices, a division of Thermo Fisher Scientific, under a contractual agreement with Duke University.

Samples in inventory: Data are summarized by sample type as well as specific container and size. Participant counts are unique individuals with one ore more aliquots. Aliquot counts are all unique samples for a given type and container, size. Freezers is a calculation of approximate storage requirements based on sample type/size, box size, and number of boxes that can be stored per freezer.

3 Participant self-reported changes in health via annual follow up. Participants are asked to complete a follow-up form once a year around the time of their original enrollment date. Participants may update contact information, primary care physician/practice and alternate contact. PROMIS domains are repeated at each annual time point in order to capture changes in participant-reported outcomes over time. The form collects new incidence/diagnosis of the same 34 medical conditions surveyed at baseline. Hospitalizations during the past year are collected along with reason, as well as specific medical procedures. Participants may update their medication list to reflect current medications, vitamins and supplements being taken at the time of follow up form completion.

Vital status: Death reported by family member or alternate contact is confirmed by obituary as the primary source. Cause of death is not captured. Follow-up metrics: Follow-up is defined as complete if participant fills out the survey online or by mail or phone. Completeness is measured as surveys completed relative to years eligible to complete follow-up. Medical conditions: "Please indicate if you have received a new diagnosis of any of the following medical conditions in the past year (yes, no, don't know)". Counts and percentages are unique participants reporting yes to specific condition in follow-up for participants that did NOT report yes at baseline. Procedures: "Please indicate if you have any of the following medical procedures in the past year". Counts are unique participants reporting the specified procedure one or more times during follow up. Hospitalizations: Participants are asked to report if they have been hospitalized within the last year, for each hospitalization they are asked to list reason(s) for hospitalization, admission date and hospital name. Reasons for hospitalization are captured as free-text responses as written by participants. Responses are coded, when possible, in order to list the most frequently reported reasons for hospitalization. Medications: (see note above for medications reported at baseline). The denominator for data based on last follow-up are participants with at least one follow-up survey complete.

4 Electronic health record (EHR) data from regional healthcare providers. Duke has partnered with regional healthcare providers to integrate data from EHR systems for consented MURDOCK Study participants. Participants are identified in EHR systems with robust matching algorithms using common identifiers from the MURDOCK and EHR databases. Data are transferred under a data use agreement (DUA) with the specific provider organization which specifies the scope of data and frequency of transfers. Data availability vary by participant and depend on whether or not a participant has had one or more encounters with the healthcare provider system during the time period included in the dataset.

Available EHR datasets: Data are summarized by healthcare provider organizations. Counts are unique participants with one or more ICD codes in the EHR dataset. Available EHR domains: Data area summarized by domain in the EHR dataset. Counts are unique participants with one of more records (rows of data) for the specified domain. Insights from available EHR data: Specific EHR data related to the population of research interest is presented with granularity when possible.

5 Additional data collection from studies with MURDOCK participants. MURDOCK Study participants may be recruited to enroll in additional research study opportunities by Duke researchers or other collaborators. Data sharing is a condition of collaboration with with the MURDOCK Study; therefore, data collected from MURDOCK Study participants and/or generated from biospecimens as part of additional research studies is returned for integration with all other MURDOCK registry data

"Storefronts" for nested sub-cohorts summarize surveys, assessments and/or other data collected specifically as part of enrollment and participation in the study. Samples in inventory: Samples are summarized if collected (see note above for samples collected at baseline). Participation in other studies: Counts are participants from the population of research interest enrolled in the specified study listed. Brief descriptions of relevant studies are listed along with a summary of study procedures and/or data collected.



Kidney disease

Implantable cardiac defibrillator

Memory & Cognitive Health Study (MHS), MURDOCK Study nested sub-cohort, N=1,595

Participant self-reported characteristics at MURDOCK Study enrollment (baseline, [February 2009 - June 2016])

Participant self-reported chara	cteristics at MURDOCK S	Study enrollmer	nt (baseline, [Februa	ary 2009	June 2016])	
Demographics at baseline		Education at	baseline				
Age	Baseline	Less than high school graduate				60 (4%)	
Median (25th, 75th)	65 (60, 71)	High school graduate, equivalent				345 (22%)	
Min, Max	49, 90+	Some college or associates degree				631 (40%)	
Sex		Bachelor's degree				317 (20%)	
Female	1,048 (66%)	Master's or higher professional degree				242 (15%)	
Male	547 (34%)	Income at baseline					
Race		Under \$10,00				48 (3%)	
American Indian & Alaska Native	2 (<1%)	\$10,000-29,999				289 (18%)	
Asian	1 (<1%)	\$30,000-49,999				347 (22%)	
Black or African American	115 (7%)	\$50,000-49,999				300 (19%)	
Native Hawaiian & Other Pacific Islander	2 (<1%)	\$70,000-89,999				195 (12%)	
White/Caucasian	1,444 (91%)	\$90,000 or more			294 (18%)		
Other	2 (<1%)	Don't know, no response				122 (8%)	
Multiple	20 (1%)					122 (070)	
Don't know/Not sure/Not answered	9 (<1%)		ndex (BMI) at baseli	ine			
Ethnicity		<18.5				11 (1%)	
Hispanic or Latino	12 (<1%)	18.5-24.9				406 (26%)	
Non-Hispanic or Latino	1,561 (98%)	25-29.9				612 (38%)	
Don't know/Not sure/Not answered	22 (1%)	30+				561 (35%)	
Smoking history at baseline		Exercise at b	paseline				
Smoked	772 (48%)	Little to no physical activity				588 (37%)	
Never smoked	813 (51%)	Weekend light exercise				219 (14%)	
Don't know, no response	10 (1%)	Moderate activity 3x per week				548 (34%)	
Current or prior medical conditions reporte		Heavy activity	3x per week			127 (8%)	
25 of 34 solicited medical conditions, listed by		Heavy activity	5x per week			103 (6%)	
High cholesterol	897 (56%)	Medications,	vitamins, supplement	ents at bas	eline		
High blood pressure	819 (51%)	Median (25th, 75th) reported				8 (5, 12)	
Osteoarthritis	498 (31%)	10+ reported, n (%)				633 (40%)	
Obesity	466 (29%)	Top 5 reported medications					
Depression	367 (23%)	Simvastatin				299 (19%)	
Skin cancer, not melanoma	329 (21%)	Lisinopril				291 (18%)	
Osteoporosis/Osteopenia	321 (20%)	Levothyroxine				281 (18%)	
Diabetes	297 (19%)	·				273 (17%)	
Thyroid disease	291 (18%)	Cholecalciferol Omeprazole				271 (17%)	
Asthma	192 (12%)						
Coronary artery disease	184 (12%)	•	rently in inventory (•			
Heart attack or angina	162 (10%)	Sample	Container, Size			Freezers	
Atrial fibrillation	150 (9%)	Plasma	Cryovial, 0.5 mL	1,353	9,071	0.160	
Gout	118 (7%)	Serum	Cryovial, 0.5 mL	1,339	5,418	0.096	
Rheumatoid arthritis	116 (7%)		Cryovial, 5.0 mL	1,433	1,433	0.051	
Other autoimmune disease	100 (6%)	Whole blood	PAXgene RNA	1,174	1,745	0.102	
Emphysema or "COPD"	86 (5%)	I lain a	Vacutainer, 2.0 mL		40	0.001	
Breast cancer	73 (5%)	Urine	Cryovial, 0.5 mL	2	2	0.000	
Melanoma	71 (4%)	DNIA	Cryovial, 10.0 mL	1,364	1,364	0.108	
Stroke	71 (4%)	DNA	Cryovial, 1.0 mL	994	996	0.015	
Other type of cancer	67 (4%)	Total			20,126	0.533	
Congestive heart failure	57 (4%)						
Prostate cancer	54 (3%)						

45 (3%)

42 (3%)

234 (15%)



Hemoglobin A1c

	Participant status	and data	from MURI	OOCK Study	y follow-up surveys and electronic health r	ecords		
Participa	nt vital status				New medical condition diagnoses reported		•	
Alive			1,2	286 (81%)	16 of 34 solicited medical conditions, listed			
Decease	d			309 (19%)	Osteoarthritis		322 / 1,097 (29%	
۸۵۵				Current	Osteoporosis/Osteopenia		1,274 (20%	
Age	OEth ZEth)				Skin cancer, not melanoma	256 /	1,266 (20%	
	25 th , 75 th)			77 (72,82)	High blood pressure	243	243 / 776 (31%	
Min, Max				64, 90+	High cholesterol	220	/ 698 (32%	
Follow-u	p metrics, study participati	on			Rheumatoid arthritis 191		1,479 (13%	
Median (25th, 75th) months since enrol	lment	153 (14	4.25, 171)	Atrial fibrillation		1,445 (11%	
Median (25th, 75th) years since enrollm	ent	,	3 (12, 15)	Thyroid disease		138 / 1,304 (11%	
Median (25th, 75th) yearly follow-ups c	omplete		10 (6, 12)	Diabetes	135 /	135 / 1,298 (10%	
	completeness of follow-up, n/N		13,538/16,4	` ' '	Coronary artery disease	132	132 / 1,411 (9%	
	one (1) follow-up survey comp	. ,		547 (97%)	Obesity	131 /	1 / 1,129 (12%	
	mpletion (n, %)			729 (46%)	Emphysema or "COPD"		124 / 1,509 (8%	
	ipleted follow-up ≤ 18 months			786 (49%)	Depression	123 /	123 / 1,228 (10%	
	in one or more other studies			95 (100%)	Congestive heart failure		/ 1,538 (7%	
Lillolled	in one of more other studies		1,00	93 (10076)	Kidney disease		100 / 1,550 (6%	
Available	e EHR datasets by source (any ICD c	ode)		Asthma		/ 1,403 (7%	
Any sour	ce		-	743 (47%)				
Novant H	Health		į	561 (35%)	Participants reporting procedures in follo	ow up		
Cabarrus	Health Alliance			247 (15%)	CT or MRI scan		1,256 (79%	
Cabarrus	Rowan Community Health C	Centers		21 (1%)	Joint x-ray		1,084 (68%	
Bethesda	a Health Center			0	Chest x-ray		1,072 (67%	
Commun	ity Free Clinic			2 (<1%)	Heart/cardiac stress test		701 (44%	
	Carolinas Healthcare)			0	Joint replacement		351 (22%	
, m. m. (c	Jaromiao Froattiroaro,				Heart/cardiac catheterization		259 (16%	
	e EHR data domains				Heart/cardiac angioplasty or stent		148 (9%	
Diagnosis	3			743 (47%)	Coronary artery bypass surgery		70 (4%	
Labs Vitals				593 (37%) 562 (35%)	Hospitalizations reported in follow up			
Medication	nns			576 (36%)	Participants reporting 1 or more hospitalizations		941 (59%)	
Allergies)			344 (22%)	Unique hospitalizations reported		1,670	
Immuniza	ations			286 (18%)	Median (25 th , 75 th) hospitalizations reported		2 (1, 3	
Problems				472 (30%)	Coded reasons for self-reported hospitalizat	ion	_ (., o	
Procedur	es		(361 (23%)	listed in descending frequency		Participant	
Hospitaliz	zations		2	274 (17%)	Uncoded	1,115	58	
Insights	from available EHR data				Surgery	238	183	
Date ran	ge: Jul. 1993 (first encounter)	, Aug. 202	2 (last encou	unter)	Knee replacement	188	130	
	of days between first and last	encounter			Hip replacement	99	7-	
,	25 th , 75 th)		1,970 (20	2, 3203.5)	Fracture		7:	
Min, Max			-1	0, 10184	Body mass index (BMI) at most recent co	85		
Phecode	hecodes, mapped from diagonal Description	Group	aes	n, ppts	` '	impleted folic	•	
272.1	Hyperlipidemia		/metabolic	11, <i>ρρι</i> ς 211	<18.5		34 (2%)	
401.1	Essential hypertension	circulatory		202	18.5-24.9		457 (30%)	
250.2	Type 2 diabetes		/metabolic	85	25-29.9		595 (38%)	
244.4	Hypothyroidism NOS	endocrine	/metabolic	77			461 (30%	
530.1	Esophagitis, GERD and related diseases	Digestive		73	Medications, vitamins, supplements at most re Median (25th, 75th) reported		•	
261.4	Vitamin D deficiency	endocrine	/metabolic	69			8 (5, 12)	
	aboratory tests				10+ reported, n (%)		540 (34%	
Test				articipants				
	lensive metabolic panel		2,451	354	Atorvastatin		42 (21%)	
	l differential etabolic panel			330 289	Levothyroxine		329 (21%)	
Lipid par	· ·		1,513	289	0		279 (17%	
TSH			1,477	279	Amlodipine		244 (15%	
	hin A1c		1,500	256	· ·		(

1,509

256

Lisinopril



Memory & Cognitive Health Study (MHS), cohort-specific visits, assessments, samples

MHS Visit 1, N=1,596

(November 2011 - July 2016)

MHS Visit 1 Assessments

Handedness questionnaire

Memory testing history

Montreal cognitive assessment (MoCA), version 7.1

ADCS - Cognitive function screen

Word list memory task, recall, recognition

Reitan trail making test - Part B

MHS Visit 2, N=880

Visit time points two years following Visit 1 (September 2014 - June 2016)

MHS Visit 2 Assessments

Handedness questionnaire

Memory testing history

Montreal cognitive assessment (MoCA), version 7.1

ADCS - Cognitive function screen

Word list memory task, recall, recognition

Reitan trail making test - Part B

Specimens in inventory, collected at Visit 2							
Sample	Container, Size	Participants	Aliquots	Freezers			
Plasma	Cryovial, 0.5 mL	697	1,485	0.026			
Serum	Cryovial, 0.5 mL	501	959	0.017			
Whole blood	Vacutainer, 3.0 mL	1	1	0.000			
	Vacutainer, 6.0 mL	1	1	0.000			
Total			2,446	0.043			