

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).

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

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. 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*.



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MURDOCK Study participants with cardiovascular disease, N=2,875

Participant self-reported characteristics at MURDOCK Study enrollment (baseline, [February 2009 - February 2018])

CVD Phenotypes in the MURD	OCK Study	
Atrial fibrillation		1,109
Heart failure		643
Peripheral arterial disease	83	
Stroke		714
Demographics at baseline		
Age		Baseline
Median (25 th , 75 th)		65 (56, 73)
Min, Max		<18, 90+
Sex		
Female		1,541 (54%)
Male		1,334 (46%)
Race		
American Indian & Alaska Native		10 (<1%)
Asian		5 (<1%)
Black or African American		331 (12%)
Native Hawaiian & Other Pacific	Islander	1 (<1%)
White/Caucasian		2,387 (83%)
Other		70 (2%)
Multiple		57 (2%)
Don't know/Not sure/Not answere	ed	14 (<1%)
Ethnicity		
Hispanic or Latino		112 (4%)
Non-Hispanic or Latino		2,694 (94%)
Don't know/Not sure/Not answere	ed	47 (2%)
Smoking history at baseline		
Smoked		1,559 (54%)
Never smoked		1,293 (45%)
Don't know, no response		23 (1%)
Current or prior medical condit 20 of 34 solicited medical conditi	tions reported ons, listed by de	at baseline escending frequency
High blood pressure		1,751 (61%)
High cholesterol		1,714 (60%)
Obesity		900 (31%)
Osteoarthritis		835 (29%)
Depression		795 (28%)
Diabetes		763 (27%)
Coronary artery disease		711 (25%)
Heart attack or angina		689 (24%)
Skin cancer, not melanoma		559 (19%)
Atrial fibrillation		542 (19%)
Thyroid disease		472 (16%)
Osteoporosis/Osteopenia		447 (16%)
Asthma		428 (15%)
Stroke		360 (13%)
Rheumatoid arthritis		330 (11%)
Congestive heart failure		291 (10%)
Emphysema or "COPD"		291 (10%)
Gout		276 (10%)
Other autoimmune disease		173 (6%)
Multiple sclerosis		156 (5%)

Education at I	baseline					
Less than high	257 (9%)					
High school graduate, equivalent			704 (24%)			
Some college	or associates degree	9	1,087 (38%)			
Bachelor's deg	Iree			499 (17%)		
Master's or hig	her professional deg	gree		324 (11%)		
Income at bas	seline					
Under \$10,000)			196 (7%)		
\$10,000-29,99	9			641 (22%)		
\$30,000-49,99	9			542 (19%)		
\$50,000-69,99	9			448 (16%)		
\$70,000-89,99	9			285 (10%)		
\$90,000 or mo	re			458 (16%)		
Don't know, no	response			305 (10%)		
Body mass in	dex (BMI) at baseli	ne				
<18.5 (underw	eight)			33 (1%)		
18.5 - 24.9 (no	rmal weight)			662 (23%)		
25 - 29.9 (over	weight)			1,036 (36%)		
30+ (obese)				1,135 (40%)		
Exercise at ba	aseline					
Little to no phy	sical activity		1,315 (46%)			
Weekend light	exercise		389 (14%)			
Moderate activ	rity 3x per week		784 (27%)			
Heavy activity	3x per week			211 (7%)		
Heavy activity	5x per week			154 (5%)		
Medications,	vitamins, suppleme	ents at basel	ine			
Median (25 th , 7	75 th) reported			9 (5, 12)		
10+ reported,	n (%)			1,218 (42%)		
Top 5 reporte	d medications (cod	led)				
Lisinopril			659 (23%)			
Simvastatin			535 (19%)			
Metoprolol			529 (19%)			
Omeprazole			520 (18%)			
Hydrochlorothi	azide		461 (16%)			
Samples currently in inventory (collected at baseline time point)						
Sample	Container, Size	Participants	Aliquots	Freezers		
Plasma	Cryovial, 0.5 mL	2,660	31,246	0.551		
Serum	Cryovial, 0.5 mL	2,678	21,689	0.383		
	Cryovial, 5.0 mL	2,378	2,379	0.084		
Whole blood	PAXgene RNA	2,508	5,282	0.308		
	Vacutainer, 2.0 mL	1,177	1,791	0.052		
Buffy coat	Cryovial, 2.0 mL	1	1	0.000		
Urine	Cryovial, 0.5 mL	7	7	0.000		
Cryovial, 10.0 mL 2,472			2,472	0.196		
Total			64,867	1.574		



Participant status and data from MURDOCK Study follow-up surveys and electronic health records

Participant	vital status					
Alive				2,100 (73%		
Deceased				775 (27%)		
Current Age				Curren		
- Median (25	th , 75 th)				73	(65, 80)
Min. Max	. ,					26. 90+
Follow-up	metrics. studv participati	on				,
Median (25	^h . 75 th) months since enrol	Iment		1,	41 (1	22 156)
Median (25	^h 75 th) years since enrollm	ent		12 (10 13)		
Median (25	h 75 th) annual follow-ups o	omnlet	P	7 (3 10		
Overall com	nleteness of follow-up, n/N	J (%)		17 648/24 933 (71%		
At least one	(1) follow-up survey com	loto ni	(0/_)	2 608 (01%		
100% com	letion (n. %)	nete, ii i	(70)		1 00)7 (35%)
	r = 10 months				1,00	(35%)
	$red 1010w-up \leq 10 1101111s$				1,31	10 (40%) 7 (50%)
Enrolled In (one of more other studies				1,48	97 (52%)
Available E	HR datasets by source (a	any ICE) co	de)		
Any source					1,35	51 (47%)
Novant Hea	lth				1,00	04 (35%)
Cabarrus H	ealth Alliance				41	4 (14%)
Cabarrus R	owan Community Health C	enters				93 (3%)
Bethesda H	ealth Center				1	4 (<1%)
Community	Free Clinic				1	1 (<1%)
Atrium (Care	olinas Healthcare)					C
Available E	HR data domains					
Diagnoses				1,351 (47%)		
Labs				1,079 (38%)		
Vitals					1,014 (35%)	
Medications	5				1,07	71 (37%)
Allergies					62	26 (22%)
Immunizatio	ons				49) 9 (17%)
Problems					87	70 (30%)
Procedures					68	38 (24%)
Hospitalizat	ions			540 (19%)		
Insights fr	om available EHP data				0	10 (1070)
Date range	luly 1993 (first encounter) Διια	2022) (last e	ncou	nter)
Number of	days between first and last	, Aug. encoui	nter		ncou	mory
Median (25	th 75th)		1903	(221 5	318	9 5)
Min. Max	,,	(0. 10	552	0.0	0.0)
Phecode	Description	Group	.,			n, ppts
401.1	Essential hypertension	, circula	tory	system		452
272.1	Hyperlipidemia	endocr	rine/ı	netabol	ic	446
250.2	Type 2 diabetes	endocr	rine/ı	netabol	ic	207
411.4	411.4 Coronary atherosclerosis circulatory system 175			175		
530.1	530.1 Esophagitis, GERD endocrine/metabolic 14			147		
261.4	261.4 Vitamin D deficiency endocrine/metable			netabol	ic	144
Select labo	oratory tests					
Test Lab			Labs	s Participant		icipants
Comprehensive metabolic panel 5,93			5,932	32 680		
CBC and di	CBC and differential 4,7;			639		
Basic Metal	polic Panel	2	4,456	56 595		
Lipid Panel 2,82			2,824	24 568		
Hemoglobir	n A1c	3	3,115	15 551		
TSH 2,52			2,527	539		

New medical condition diagnoses reported in follow-up 17 of 34 solicited medical conditions, listed by descending frequency					
Atrial fibrillation		538 / 1	2,333 (23%)		
Osteoarthritis		486 / 1	2,040 (24%)		
Coronary artery disease		457 / :	2,164 (21%)		
High cholesterol		365 /	1.161 (31%)		
Rheumatoid arthritis		348 /	2.545 (14%)		
Skin cancer not melanoma		328 /	2 316 (14%)		
Concestive heart failure		326 / 3	2.584 (13%)		
Stroke		322 / 1	2.515 (13%)		
Osteoporosis/Osteopenia		314 /	2 428 (13%)		
Heart attack or angina		308 /	2.186 (14%)		
High blood pressure		308 /	1 124 (27%)		
Emphysema or "COPD"		260 / 3	2 584 (10%)		
Depression		253 /	2 080 (12%)		
Diabetes		240 /	2 112 (11%)		
Thyroid disease		239/	2,112 (11%)		
Obesity		232 /	1 975 (12%)		
Kidney disease		2027	/ 2 748 (8%)		
Procedures reported in follow	n		2,740 (070)		
CT or MDI agon	μþ		0 107 (700/)		
			2,107 (73%) 1.010 (67%)		
Leist x-ray			1,919 (07%)		
Joint x-ray			1,044 (57%)		
Heart/cardiac stress lest		1,428 (50%)			
Hearl/cardiac callelenzation		727 (25%)			
Joint replacement		310 (10%) 442 (15%)			
Hearl/cardiac angioplasty or sten		443 (15%)			
Coronary artery bypass surgery			198 (7%)		
Hospitalizations reported in fol	low up				
Participants reporting 1 or more r	nospitalizations		1,719 (60%)		
Unique hospitalizations reported		4,750			
Median $(25^{\text{m}}, 75^{\text{m}})$ hospitalization	is reported		2 (1, 3)		
listed in descending frequency	nospitalization	Events	Participants		
Uncoded		2 184	1 099		
Surgery		385	202		
Knee Replacement		235	176		
Stroke		230	184		
AFIB		200	161		
Body mass index (BMI) at mos	t recent compl	eted follo	wun		
<18.5 (underweight)	r recent compr	eteu iono	40 (00()		
18.5 24.0 (pormal woight)			40 (2%)		
25 29 9 (overweight)			706 (27%)		
20 - 29.9 (Overweight)		932 (36%)			
			921 (35%)		
wedications, vitamins, suppler	nents at most	recent fol			
Integran (25 ^{url} , 75 ^{url}) reported		8 (5, 12)			
Ton Francestad and the state			984 (34%)		
Top 5 reported medications			050 (000)		
Atorvastatin			652 (23%)		
ivietoproioi			632 (22%)		
Lisinoprii			485 (17%)		
			481 (17%)		
Levothyroxine			455 (16%)		

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MURDOCK Study participants with cardiovascular disease, N=2,875

Cardiovascular disease phenotypes in the MURDOCK Study

Atrial fibrillat	ion			n=1,109
Source of dia	ignosis			
Self-report onl	У			989
Self-report & E	EHR			91
EHR only				29
Samples curr	ently in inventory (collected at b	baseline t	ime point)
Sample	Container, Size	Participants	Aliquots	Freezers
Plasma	Cryovial, 0.5 mL	1,036	11,989	0.211
Serum	Cryovial, 0.5 mL	1,032	8,155	0.144
	Cryovial, 5.0 mL	926	926	0.033
Whole blood	PAXgene RNA	974	1,970	0.115
	Vacutainer, 2.0 mL	406	616	0.018
Buffy coat	Cryovial, 2.0 mL	0	0	0.000
Urine	Cryovial, 0.5 mL	4	4	0.000
	Cryovial, 10.0 mL	952	952	0.076
Total			24,612	0.596

Heart failure				N=643		
Source of dia	gnosis					
Self-report only	y			579		
Self-report & E	HR			38		
EHR only				26		
Samples curr	ently in inventory (collected at b	baseline t	ime point)		
Sample	Container, Size	Participants	Aliquots	Freezers		
Plasma	Cryovial, 0.5 mL	595	6,955	0.123		
Serum	Cryovial, 0.5 mL	594	4,583	0.081		
	Cryovial, 5.0 mL	509	509	0.018		
Whole blood	PAXgene RNA	557	1,166	0.068		
	Vacutainer, 2.0 mL	251	379	0.011		
Buffy coat	Cryovial, 2.0 mL	0	0	0.000		
Urine	Cryovial, 10.0 mL	548	548	0.043		
Total			14,140	0.344		

Stroke				n=714	
Source of diagnosis					
Self-report onl	У			653	
Self-report & E	EHR			29	
EHR only				32	
Samples curr	ently in inventory (collected at k	baseline t	ime point)	
Sample	Container, Size	Participants	Aliquots	Freezers	
Plasma	Cryovial, 0.5 mL	656	7,576	0.134	
Serum	Cryovial, 0.5 mL	656	5,356	0.094	
	Cryovial, 5.0 mL	584	585	0.021	
Whole blood	PAXgene RNA	620	1,313	0.077	
	Vacutainer, 2.0 mL	297	434	0.013	
Buffy coat	Cryovial, 2.0 mL	1	1	0.000	
Urine	Cryovial, 0.5 mL	1	1	0.000	
	Cryovial, 10.0 mL	611	611	0.048	
Total			15,877	0.386	

Peripheral art		n=83			
Source of dia	gnosis				
Self-report onl	у			14	
Self-report & E	HR			1	
EHR only				68	
Samples curr	ently in inventory (collected at b	baseline t	ime point)	
Sample	Container, Size	Participants	Aliquots	Freezers	
Plasma	Cryovial, 0.5 mL	78	873	0.015	
Serum	Cryovial, 0.5 mL	80	620	0.011	
	Cryovial, 5.0 mL	61	61	0.002	
Whole blood	PAXgene RNA	75	157	0.009	
	Vacutainer, 2.0 mL	33	53	0.002	
Buffy coat	Cryovial, 2.0 mL	0	0	0.000	
Urine	Cryovial, 10.0 mL	73	73	0.006	
Total			1,837	0.045	