

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 phy sical exam (vital signs, height, 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 cry ovials. Urine was collected and aliquoted in cry ovials. Serial 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.*



Melanoma

MURDOCK Study participants with obese BMI classification at baseline, N=4,608

Participant self-reported characteristics at MURDOCK Study enrollment (baseline, February 2009 - March 2018

Dem ographics at baseline	Education at	baseline					
Age	Baseline			Less than high school graduate			
Median (25th, 75th)	^h , 75 th) 52 (41, 62)			High school graduate, equivalent			
Min, Max	<18, 90+	Some college	e or associates degree		1,800 (39%)		
Sex		Bachelor's de	gree			765 (17%)	
Female	ale 3,103 (67%)			Master's or higher professional degree			
Male	1,505 (33%)	Master's or higher professional degree 430 (9% Income at baseline					
Race		Under \$10,000 365 (8%)					
American Indian & Alaska Native	21 (<1%)	\$10,000-29,999			912 (20%)		
Asian	11 (<1%)	\$30,000-49,9				851 (18%	
Black or African American	869 (19%)	\$50,000-69,9			649 (149		
Native Haw aiian & Other Pacific Islander	3 (<1%)	\$70,000-89,9			462 (10%		
White/Caucasian	3,097 (67%)	\$90,000 or more			695 (15%)		
Other	453 (10%)	Don't know, no response		674 (14%)			
Multiple	101 (2%)						
Don't know /Not sure/Not answ ered	Body mass index (BMI) at baseline						
Ethnicity	53 (1%)	<18.5 (underw eight)			0		
Hispanic or Latino	594 (13%)	18.5 - 24.9 (normal w eight)			0		
Non-Hispanic or Latino	3,951 (86%)	25 - 29.9 (overweight)			0		
Don't know /Not sure/Not answ ered	63 (1%)	30+ (obese)			4	1,608 (100%	
	66 (176)	Exercise at baseline					
Smoking history at baseline Smoked	Little to no physical activity			2,178 (47%)			
	1,922 (42%)	Weekend light exercise		984 (21%)			
Never smoked	2,648 (57%)	Moderate activity 3x per w eek		1,073 (23%			
Oon't know, no response 38 (1%)		Heavy activity 3x per w eek			222 (5%)		
Current or prior medical conditions reported a 25 of 34 solicited medical conditions, listed by de	Heavy activity	Heavy activity 5x per w eek 124 (3%)					
Obesity	2,722 (59%)	Medications	s, vitamins, supplen	nents at base	line		
High blood pressure	2,293 (50%)	Median (25th,	75th) reported		5 (2, 10		
High cholesterol	2,057 (45%)	10+ reported, n (%)			1,157 (25%)		
Depression	1,376 (30%)	Top 5 reported medications				, . (
Diabetes	1,132 (25%)					792 (17%	
Osteoarthritis	953 (21%)	Hydrochlorothiazide			718 (16%)		
Asthma	787 (17%)	•					
Thyroid disease	657 (14%)	Metformin			708 (15%)		
Rheumatoid arthritis	433 (9%)	Levothyroxine			578 (13%)		
Multiple sclerosis	340 (7%)	Omeprazole				552 (12%	
Skin cancer, not melanoma	335 (7%)		rently in inventory				
Osteoporosis/Osteopenia	331 (7%)	Sam ple	Container, Size	Participants	Aliquots	Freezers	
Heart attack or angina	287 (6%)	Plasma	Cryovial, 0.5 mL	4,279	52,728	0.930	
Coronary artery disease	279 (6%)	Serum	Cryovial, 0.5 mL	4,329	38,758	0.684	
Gout	279 (6%)		Cryovial, 5.0 mL	3,694	3,694	0.130	
Other autoimmune disease	270 (6%) 258 (6%)	Whole blood	PAXgene RNA	4,117	9,589	0.559	
Emphysema or "COPD"	248 (5%)		Vacutainer, 2.0 mL	2,331	3,788	0.110	
Other mental illness	234 (5%)	Buffy coat	Cryovial, 2.0 mL	0	0	0.000	
Atrial fibrillation	194 (4%)	Urine	Cryovial, 4.0 mL	11	11	0.000	
Stroke	152 (3%)		Cryovial, 10.0 mL	4,152	4,152	0.330	
		Total			112,720	2.743	
Congestive heart failure	147 (3%)						
Kidney disease Other type of capeer	132 (3%)						
Other type of cancer	132 (3%)						
Breast cancer	100 (2%)						

95 (2%)



MURDOCK Study participants with obese BMI classification at baseline, N=4,608

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

		us and da	ta from MURI	OOCK Stud	y follow-up surveys and electronic health recor				
Participa: Alive	nt vital status		<i>A</i> (090 (89%)	New medical condition diagnoses reported in follow-up 17 of 34 solicited medical conditions, listed by descending frequency				
Deceased				518 (11%)	Osteoarthritis	607 /	3,655 (17%)		
Current A			,	Current	High blood pressure	537 / 2,315 (23%)			
	_		6		High cholesterol	igh cholesterol 534 / 2,551			
Median (2	oui, 7 oui)		0	31 (51, 72)	Obesity	463 /	1,886 (25%)		
Min, Max	n matrice etudy participa	ation		26, 90+	Diabetes	454 / 3,476 (13%)			
	Follow-up metrics, study participation			(100 150)	Rheumatoid arthritis	408 / 4,175 (10%)			
,	5th, 75th) months since enro			(122, 158)	Depression	3,232 (11%)			
,	5 th , 75 th) years since enroll			2 (10, 13)	Osteoporosis/Osteopenia				
	5th, 75th) annual follow -ups			5 (1, 9)	Thyroid disease	293 / 3,951 (7%)			
	mpleteness of follow -up, n	` '	24,628/43,2		Skin cancer, not melanoma	278 / 4,273 (7%)			
At least or	ne (1) follow -up survey cor	nplete, n (%			Other autoimmune disease				
100% com	pletion (n, %)		1,0	074 (23%)	Atrial fibrillation				
Last comp	leted follow-up≤18 month	IS	1,9	999 (43%)		221 / 4,414 (5% 205 / 4,476 (5%			
Enrolled in	one or more other studies	3	1,8	376 (41%)	Kidney disease				
Available	EHR datasets by source	(any ICD	code)		Asthma		/ 3,821 (5%)		
Any sourc	any source			242 (49%)	Emphysema or "COPD"	193 / 4,360 (4%			
Novant He	alth		1,4	487 (32%)	Gout	193 / 4,338 (4%)			
Cabarrus H	Health Alliance			343 (18%)	Coronary artery disease 176 / 4,32				
Cabarrus F	us Row an Community Health Centers 305 (7%)			· ·	Procedures reported in follow up				
	Health Center			58 (1%)	CT or MRI scan		2,585 (56%)		
	Free Clinic			42 (1%)	Joint x-ray		2,083 (45%)		
,	rolinas Healthcare)			72 (170)	Chest x-ray		2,048 (44%)		
`	EHR data domains				Heart/cardiac stress test		1,176 (26%)		
			2.0	242 (400/)	Joint replacement		622 (13%)		
Diagnoses)			242 (49%)	Heart/cardiac catheterization	392 (9%)			
Labs				333 (40%)	Heart/cardiac angioplasty or stent	202 (4%			
Vitals					Coronary artery bypass surgery	116 (3%)			
Medication	ns .			767 (38%)	Hospitalizations reported in follow up				
Allergies	ů ,			341 (18%) 389 (15%)	Participants reporting 1 or more hospitalizations		1,820 (39%)		
lmmunizati	ons				Unique hospitalizations reported	2,949			
Problems				188 (26%)	Median (25th, 75th) hospitalizations reported	2 (1, 3)			
Procedure Hospitaliza	-			929 (20%) 774 (17%)	Coded reasons for self-reported hospitalization listed in descending frequency	Events	Participants		
Insiahtsf	rom available EHR data			, ,	Uncoded	2,070	1,127		
_	e: June 1993 (first encount	ter), Aug. 2	2022 (last enc	ounter)	Surgery	460	366		
_	f days between first and las		er:		Knee Replacement	339	241		
Median (2	5 th , 75 th)		1,851 (2	242, 3343)					
Min, Max		_	_	0, 10563	Pneumonia	136	96		
-	ecodes, mapped from di		odes		Chest pain	133	115		
Phecode 401.1	Description Essential hypertension	Group	ry system	<i>n, ppt</i> s 551	Body mass index (BMI) at most recent comp	letedfoll	ow up		
272.1	Hyperlipidemia		ne/metabolic	509	<18.5 (underw eight)	10 (0%)			
278.1	Obesity		ne/metabolic	339	18.5 - 24.9 (normal w eight)	115 (3%)			
250.2	Type 2 diabetes	endocrin	ne/metabolic	298	25 - 29.9 (overweight)	635 (16%)			
278.11	Morbid obesity		ne/metabolic	228	30+	3,147 (81%)			
530.11	GERD	digestive		206	Medications, vitamins, supplements at most	recent follow up			
296.2	Depression	mental d	isorders	180	Median (25th, 75th) reported		6 (3, 10)		
Test	boratory tests		Loho Do	rtiainanta	10+ reported, n (%)		1,015 (22%)		
	nsive metabolic panel		6,884	articipants 994	Top 5 reported medications				
	nd differential		5,529	922	Metformin	675 (15%)			
Hemoglobi			4,285	784	Lisinopril		652 (14%)		
TSH			3,106	729	Atorvastatin	609 (13%)			
	Lipid panel		3,141	710	Levothyroxine	586 (13%)			
	abolic panel		3,583	698	Omeprazole	560 (12%)			
CBC		3,114	630	Onepiazole		JUU (12%)			