



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 (CTS).

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. Serial sample collection was not done systematically for MURDOCK enrollees; however, some nested subcohorts 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 currently 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 or 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 or 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 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.*

**MURDOCK Study participants with lung cancer, N=165**
**Participant self-reported characteristics at MURDOCK Study enrollment (baseline, February 2009– February 2018)**
**Demographics at baseline**

	Baseline
<b>Age</b>	
Median (25 <sup>th</sup> , 75 <sup>th</sup> )	66 (56, 73)
Min, Max	25, 89
<b>Sex</b>	
Female	104 (63%)
Male	61 (37%)
<b>Race</b>	
American Indian & Alaska Native	0
Asian	0
Black or African American	15 (9%)
Native Hawaiian & Other Pacific Islander	0
White/Caucasian	142 (86%)
Other	5 (3%)
Multiple	3 (2%)
Don't know /Not sure/Not answered	0
<b>Ethnicity</b>	
Hispanic or Latino	7 (4%)
Non-Hispanic or Latino	153 (93%)
Don't know /Not sure/Not answered	5 (3%)
<b>Smoking history at baseline</b>	
Smoked	112 (68%)
Never smoked	50 (30%)
Don't know , no response	3 (2%)

**Current or prior medical conditions reported at baseline**
*26 of 34 solicited medical conditions, listed by descending frequency*

High cholesterol	98 (59%)
High blood pressure	80 (48%)
Lung cancer	45 (27%)
Osteoarthritis	42 (25%)
Depression	39 (24%)
Emphysema or "COPD"	38 (23%)
Obesity	37 (22%)
Skin cancer, not melanoma	32 (19%)
Osteoporosis/Osteopenia	31 (19%)
Diabetes	29 (18%)
Asthma	25 (15%)
Rheumatoid arthritis	25 (15%)
Thyroid disease	25 (15%)
Coronary artery disease	20 (12%)
Atrial fibrillation	19 (12%)
Other type of cancer	19 (12%)
Heart attack or angina	17 (10%)
Stroke	15 (9%)
Breast cancer	14 (8%)
Melanoma	12 (7%)
Congestive heart failure	10 (6%)
Prostate cancer	9 (5%)
Other autoimmune disease	8 (5%)
Multiple sclerosis	7 (4%)
Implantable cardiac defibrillator	6 (4%)
Other mental illness	6 (4%)

**Education at baseline**

Less than high school graduate	12 (7%)
High school graduate, equivalent	43 (26%)
Some college or associates degree	61 (37%)
Bachelor's degree	25 (15%)
Master's or higher professional degree	24 (15%)

**Income at baseline**

Under \$10,000	8 (5%)
\$10,000-29,999	35 (21%)
\$30,000-49,999	37 (22%)
\$50,000-69,999	25 (15%)
\$70,000-89,999	17 (10%)
\$90,000 or more	29 (18%)
Don't know , no response	14 (9%)

**Body mass index (BMI) at baseline**

<18.5 (under weight)	3 (2%)
18.5 - 24.9 (normal weight)	46 (28%)
25 - 29.9 (overweight)	68 (41%)
30+ (obese)	48 (29%)

**Exercise at baseline**

Little to no physical activity	67 (41%)
Weekend light exercise	28 (17%)
Moderate activity 3x per week	48 (29%)
Heavy activity 3x per week	14 (8%)
Heavy activity 5x per week	6 (4%)

**Medications, vitamins, supplements at baseline**

Median (25 <sup>th</sup> , 75 <sup>th</sup> ) reported	8 (5, 12)
10+ reported, n (%)	64 (39%)

**Top 5 reported medications**

Omeprazole	35 (21%)
Lisinopril	32 (19%)
Simvastatin	28 (17%)
Albuterol	25 (15%)
Hydrochlorothiazide	25 (15%)

**Samples currently in inventory (collected at baseline time point)**

Sample	Container, Size	Participants	Aliquots	Freezers
Plasma	Cryovial, 0.5 mL	149	1,733	0.031
Serum	Cryovial, 0.5 mL	156	1,214	0.021
	Cryovial, 5.0 mL	140	140	0.005
Whole blood	PAXgene RNA	143	291	0.017
	Vacutainer, 2.0 mL	55	85	0.002
Buffy coat	Cryovial, 2.0 mL	0	0	0.000
Urine	Cryovial, 10.0 mL	144	144	0.011
Total			3,607	0.087

## MURDOCK Study participants with lung cancer, N=165

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

Participant vital status	
Alive	95 (58%)
Deceased	70 (42%)
Current Age	
Median (25 <sup>th</sup> , 75 <sup>th</sup> )	73 (64, 81)
Min, Max	36, 90+

Follow-up metrics, study participation	
Median (25 <sup>th</sup> , 75 <sup>th</sup> ) months since enrollment	150 (128, 163)
Median (25 <sup>th</sup> , 75 <sup>th</sup> ) years since enrollment	12 (11, 13)
Median (25 <sup>th</sup> , 75 <sup>th</sup> ) yearly follow -ups complete	7 (4, 10)
Overall completeness of follow-up, n/N (%)	1,133 / 1,430 (79%)
At least one (1) follow -up survey complete, n (%)	160 (97%)
100% completion (n, %)	75 (45%)
Last completed follow -up ≤ 18 months	70 (42%)
Enrolled in one or more other studies	95 (58%)

Available EHR datasets by source (any ICDcode)	
Any source	67 (41%)
Novant Health	56 (34%)
Cabarrus Health Alliance	11 (7%)
Cabarrus Row an Community Health Centers	5 (3%)
Bethesda Health Center	0
Community Free Clinic	0
Atrium (Carolinas Healthcare)	0

Available EHR data domains	
Diagnoses	67 (41%)
Labs	59 (36%)
Vitals	55 (33%)
Medications	58 (35%)
Allergies	43 (26%)
Immunizations	36 (22%)
Problems	49 (30%)
Procedures	39 (24%)
Hospitalizations	32 (19%)

Insights from available EHR data	
Date range: April 1996 (first encounter), Aug. 2022 (last encounter)	
Number of days between first and last encounter:	
Median (25 <sup>th</sup> , 75 <sup>th</sup> )	2,344 (878.5, 3137.5)
Min, Max	0, 7678

Select phecodes, mapped from diagnosis codes			
Phecode	Description	Group	n, ppts
272.1	Hyperlipidemia	endocrine/metabolic	29
401.1	Essential hypertension	circulatory system	26
530.11	GERD	Digestive	14
165.1	Cancer of bronchus; lung	Neoplasms	13
250.2	Type 2 diabetes	endocrine/metabolic	13
512.7	Shortness of breath	Respiratory	11

Select laboratory tests		
Test	Labs	Participants
Comprehensive metabolic panel	429	43
CBC and differential	383	40
Lipid panel	190	35
TSH	173	35
Hemoglobin A1C	195	34
Basic metabolic panel	243	32
CBC	137	31

New medical condition diagnoses reported in follow-up	
15 of 34 solicited medical conditions, listed by descending frequency	
Lung cancer	113 / 120 (94%)
Emphysema or "COPD"	31 / 127 (24%)
High blood pressure	28 / 85 (33%)
Osteoarthritis	25 / 123 (20%)
Osteoporosis/Osteopenia	21 / 134 (16%)
Skin cancer, not melanoma	21 / 133 (16%)
Depression	21 / 126 (17%)
Thyroid disease	20 / 140 (14%)
High cholesterol	20 / 67 (30%)
Atrial fibrillation	19 / 146 (13%)
Congestive heart failure	18 / 155 (12%)
Other type of cancer	18 / 146 (12%)
Coronary artery disease	17 / 145 (12%)
Asthma	16 / 140 (11%)
Rheumatoid arthritis	16 / 140 (11%)

Procedures reported in follow up	
CT or MRI scan	144 (87%)
Chest x-ray	132 (80%)
Joint x-ray	84 (51%)
Heart/cardiac stress test	63 (38%)
Heart/cardiac catheterization	27 (16%)
Joint replacement	20 (12%)
Heart/cardiac angioplasty or stent	15 (9%)
Coronary artery bypass surgery	9 (5%)

Hospitalizations reported in follow up		
Participants reporting 1 or more hospitalizations	100 (61%)	
Unique hospitalizations reported	182	
Median (25 <sup>th</sup> , 75 <sup>th</sup> ) hospitalizations reported	2 (1, 3)	
Coded reasons for self-reported hospitalization listed in descending frequency	Events	Participants
Uncoded	121	57
Cancer	26	24
Surgery	24	16
Pneumonia	12	11
Stroke	11	7
AFIB	10	7

Body mass index (BMI) at most recent completed follow up	
<18.5 (underweight)	11 (7%)
18.5 - 24.9 (normal weight)	61 (38%)
25 - 29.9 (overweight)	51 (32%)
30+	37 (23%)

Medications, vitamins, supplements at most recent follow up	
Median (25 <sup>th</sup> , 75 <sup>th</sup> ) reported	8 (4, 12)
10+ reported, n (%)	56 (34%)

Top 5 reported medications	
Atorvastatin	36 (22%)
Omeprazole	32 (19%)
Cholecalciferol	31 (19%)
Levothyroxine	31 (19%)
Lisinopril	28 (17%)