

Cohort Overview

Mia M. Gaudet, Ph.D.

Connect Chief Scientist

Trans Divisional Research Program
Division of Cancer Epidemiology and Genetics
National Cancer Institute



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Study Objectives

Comprehensive research resource using new technologies and methods for the scientific community to study:

Cancer Etiology

Precursor to Tumor Progression Cancer Risk Prediction Early Detection of Cancer Survivorship



Connect Designed to Address Key Research Priorities



Emerging exposures



Novel biomarkers, genomics



Cutting-edge methodology



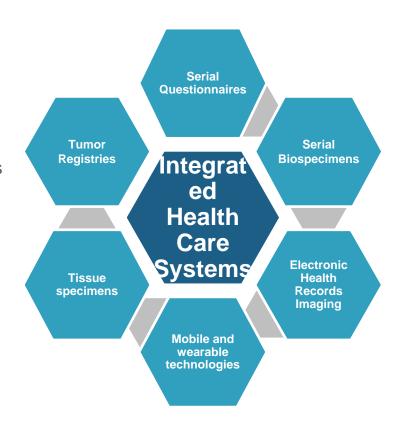
Diverse and special populations

Cohort Study Design Overview & Recruitment Update



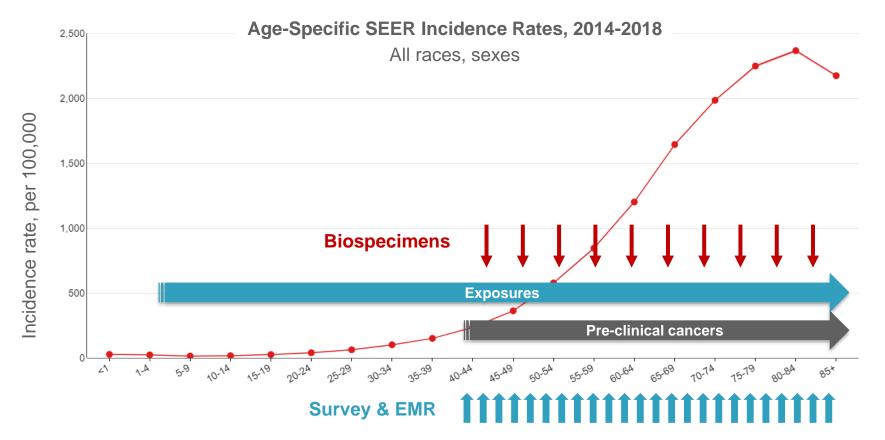
Key Study Design Features of Connect

- ► 200,000 adults across the US
 - ✓ Aged 40-65 years Expanding to 30-70 in Jan 2024
 - ✓ No history of cancer
 - √ Patients or members of partner health care systems
- ► Long-term follow-up
- Serial assessments
- Comprehensive cancer and precancer outcomes
- ► Flexible infrastructure for enhancement studies

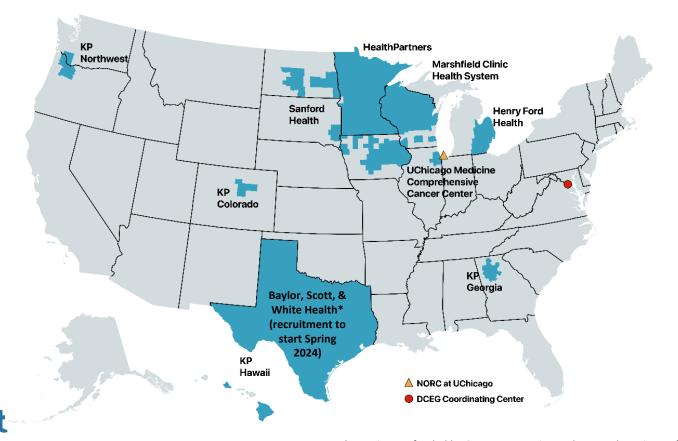




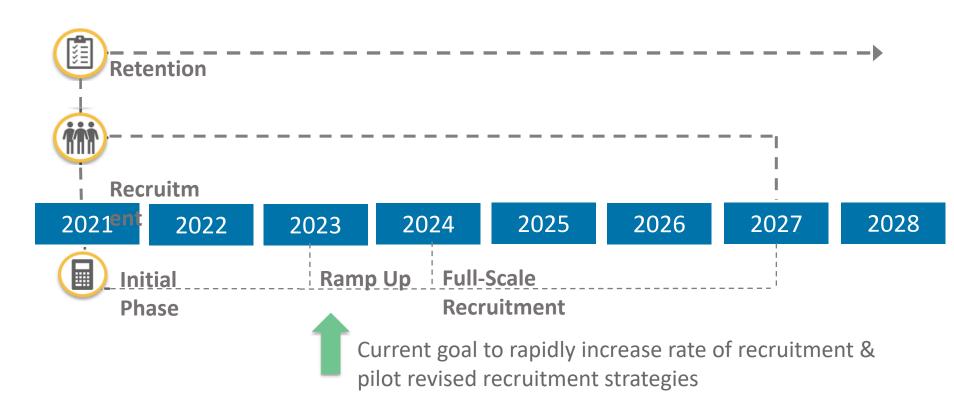
Serial Assessments to Observe Longitudinal Changes



Recruitment through 10 U.S. Integrated Health Care Systems



Scaling up recruitment



Participant Engagement

Recruitment Materials & Strategy

- Value proposition to anchor communications
- Prevention-focused messaging
- Materials designed to engender trust, enthusiasm, and inclusivity
- Monetary and non-monetary incentives
- Active and passive recruitment

Connect today. Prevent cancer tomorrow.



Retention Strategies

- Candid messaging about participant burden for the long-term
- Responsive communication using multiple channels
- Incentives for retention, such as return of results





Ongoing Engagement Strategies

- Digital engagement
- Regular contact every 6 months beyond those for study activities
- "Study anniversary" mailing
- Monitor real-time metrics to pivot engagement strategy





MyConnect Participant App



MyConnect Participant App: Communication and Study Activity Hub

Existing Functionality

Reminders & Communication







Study Activities







Planned Functionality

Return of Results

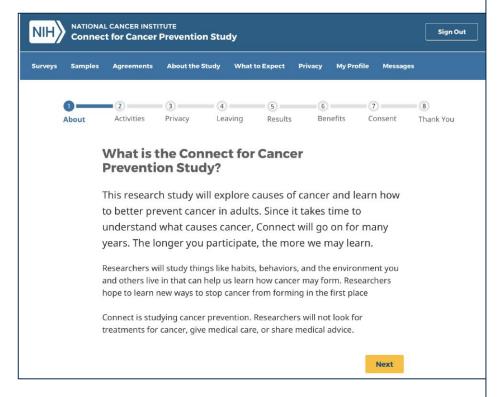


Mobile/ Wearable Technologies





Educational Module before Long Form Consent







Informed Consent

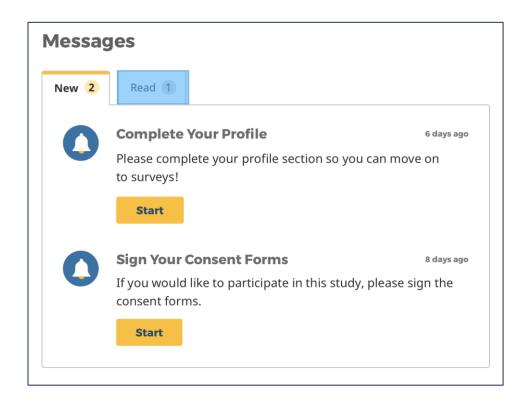
In order to join Connect, we need to you to give your informed consent by reviewing the full consent form and electronic health records release form, and signing your name.

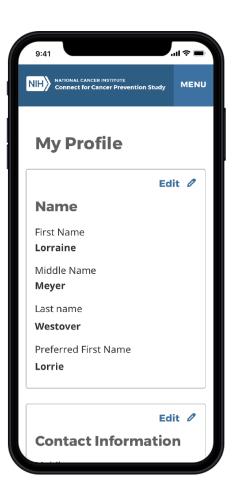
Informed consent form

This is a more detailed explanation of what it means to take part in Connect.

Download an unsigned copy of the consent form

Messages and Data Input

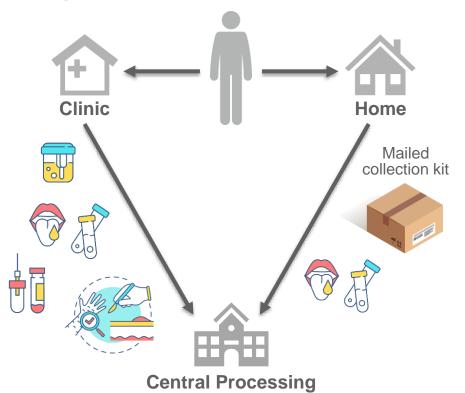




Biospecimen Collections



Biospecimen Collections at Recruitment



& Biorepository

	No. Tubes	Fraction	Tube Type
Buccal	1		Saliva-mouthwash kit
	2	Serum	10ml Red top
Dland	1	Plasma/ buffy coat	10ml Lavender top (EDTA)
Blood	1	Plasma/ buffy coat	10ml Green top (Heparin)
	1	Whole blood	10ml ACD Streck
Urine	1		10ml Urine kit
Tissue			Precursor tissue blocks, if available



Biospecimen Collections at Follow-up

- Timed on average every three years
 - More frequent for participants at higher risk of cancer
- Series of same biospecimens and new types of biospecimens (e.g., liquid biopsy, stool)
- Located at healthcare system, external clinic, home
- Paired with questionnaires and other relevant data collection



Surveys



Baseline Surveys

Background and Overall Health

- Background Information
- Medical History
- Family History of Cancer
- Education and Occupation

Medications, Reproductive Health, Exercise, Sleep

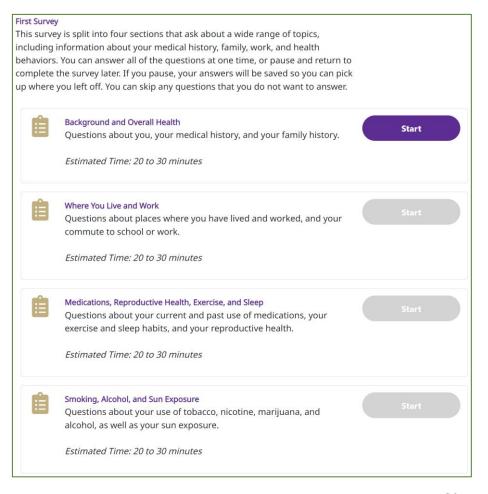
- Medications
- Pregnancy History
- Physical Activity
- Sleep

Smoking, Alcohol, Sun Exposure

- Tobacco
- Marijuana
- Alcohol
- Sun Exposure

Where You Live and Work

- Residential History
- Commuting



Surveys Accompanying Biospecimen Collection

Blood & Urine

- Recent exposures to food, drink, pharmaceuticals
- Reproductive Health
 - Last menstrual period date

Mouthwash

- Oral health and care
- Recent exposures to toothpaste, mouthwash, gum, tobacco
- Permanent teeth lost

COVID-19

- Exposure
- Complications
- Long-COVID
- Vaccination



Menstrual Cycle

Next menstrual period date

Surveys available in MyConnect

Baseline	Background & Overall Health	Demographics, medical history, family history of cancer, education & occupation	
	Medications, Reproductive Health, Exercise, Sleep	OTC/rx, menstrual cycle, pregnancy, fertility, hormonal meds & contraception, physical activity, shiftwork	
	Smoking, Alcohol, Sun Exposure	Tobacco/nicotine, marijuana, alcohol, tanning	
	Where You Live & Work	Residential, seasonal, & childhood addresses, water source, tap water intake, work & school commute	
	SSN	Full or partial SSN	
Biospecimens	Blood, Urine, Mouthwash (Research Appt)	Symptoms/behaviors on day of donation, meds/antibiotics, recent reproductive health, oral hygiene & health	
	Blood/Urine (Clinical Collection)	Symptoms/behaviors on day of donation, meds, recent reproductive health	
	COVID-19	# Infections, symptoms, hospitalization, vaccination	
	Menstrual Cycle Survey	Last period, triggered by research appt/clinical collection response	

Surveys In Development

Participant Experience	MyConnect, study communications, activities completed, overall experience, motivation for joining	
Cancer Screening History	Organ Inventory (born with/current), history of cancer screening tests	
Cancer Diagnosis Surveys (17)	Dx, symptoms, Patient-Provider interaction, medical history repeat assessment	
Menstrual & Intimate Care Products	Vaginoplasty, powder, douching, vaginal cleansing products, menstrual products	
Fecal Collection	Donation, bowel movements, meds, supplements, probiotics	
Social Determinants of Health (SDOH)	Discrimination, police interaction, medical mistrust, social support, and financial, food, and housing insecurity	
Hair Products	Dyes, relaxers, straighteners, perms, oils	
Menstrual Experience Survey	Menstrual problems, endometriosis dx and treatment	
Mothballs & Scented Products	Household exposure to p-DCB & Napthalene	

Detailed Status Available <u>Here</u>

Planned Survey Cadence

6 Months

 Diet food frequency questionnaire (DHQ)

Within 24 Months

 Repeated 24hour recalls for diet (ASA24) and physical activity (ACT24)

12 Months

- Screening History
- Quality of life (PROMIS)
- Psychosocial
- Social determinants of health

2 Years

- Baseline exposures repeat assessment
- Consumer products (Menstrual & intimate care, Hair, mothballs & scented)
- Menstrual experience (Endometriosis)

3-6 Months Post-Cancer Diagnosis Rapid Case Ascertainment

- Cancer Diagnosis Surveys (Dx, symptoms, patient-provider interaction, medical history)
 - Quality of life (PROMIS: repeat assessment & patient specific)

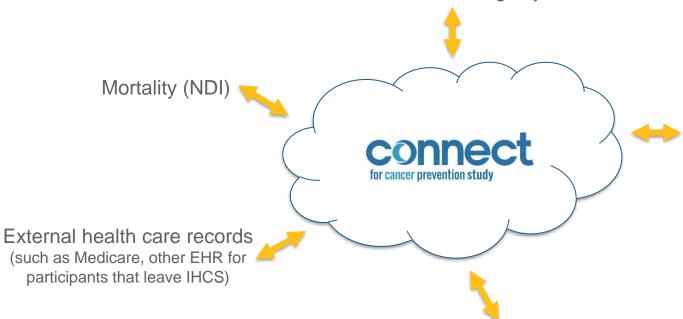
Data Linkages

Data Linkages

Mortality (NDI

(such as Medicare, other EHR for participants that leave IHCS)

Cancer registries, virtual tumor registry



Geospatial data (such as outdoor air quality data, other EPA MyEnvironment, WATERS data, US Census)

Health outcomes data (such as HIV registry)

Geographic Information

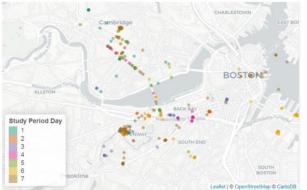
 Geocode 20-year address histories (collected on baseline survey)

 Enable linkages to datasets for numerous external risk factor and exposure information

Useful for planning ancillary field studies

Add daily movement and travel to exposure information via MyConnect

information via MyConnect









Wearable Devices and Apps

- Connect will have research-grade and consumer-oriented devices available
- Key domains where tech can improve assessment of exposures and risk factors:
 - sleep
 - light at night
 - air pollution
 - physical activity





Estimated Outcomes

Estimated total number of incident invasive cancers

Campan alta	Total Number of Expected Incident Cancers		
Cancer site	10 years of follow up	15 years of follow up	
Prostate (males only)	1,951	3,595	
Breast (females only)	1,566	2,553	
Lung & Bronchus	1,151	2,165	
Colorectal	877	1,566	
Melanoma of the Skin	410	681	
Non-Hodgkin Lymphoma	369	649	
Uterine Corpus	345	579	
Urinary Bladder	327	622	
Thyroid	232	356	
Pancreas	221	410	
Leukemia	203	365	
Ovary	146	246	

Estimated Total Number of Precursor Lesions

	Total Number of Expected Precursor Lesions		
Lesions	Retrospective at Baseline	Prospective at 10 Years of Follow Up	
Colorectal Adenomas	5,778	18,311	
Benign Breast Biopsies (females only)	2,339	4,942	
Dysplastic Nevus	2,311	3,779	
Cervical Biopsies (females only)	2,137	2,822	
High-Grade Prostatic Intraepithelial Neoplasia (males only)	238	384	
Lung Nodules	165	349	

Data System Infrastructure & Digital Tools

Designed to maximize F.A.I.R. principles



F.A.I.R. Data Ecosystem Makes F.A.I.R. Data

Principles





Accessible

 Retrievable data by identifiers based on a shared governance model



Interoperable

 Data architecture that allows for integration of data from a range of sources and applications



Reusable

 Goal of F, A, and I to allow reuse of data, code, and tools

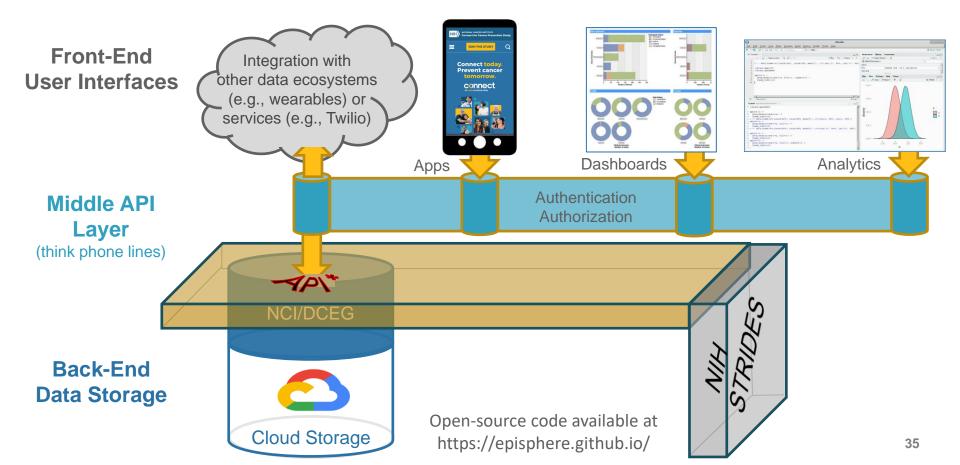
Implementation in Connect

- Online, searchable data dictionary
- Map variables to common thesaurus to aid pooling (planned)

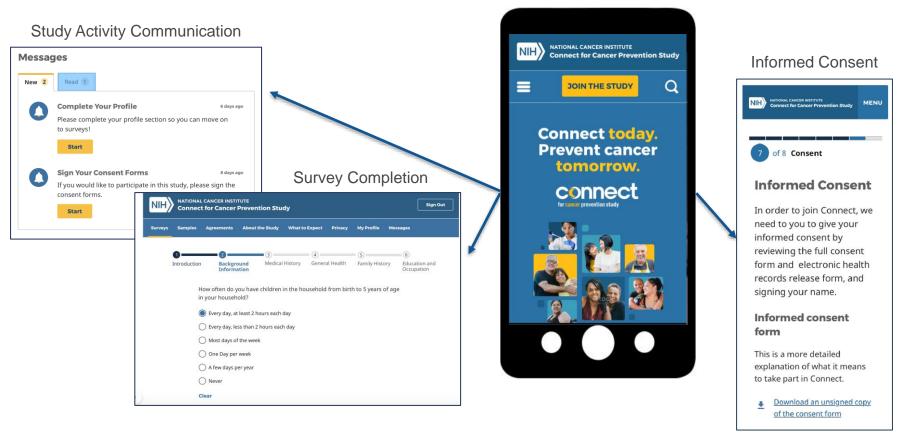
- Authentication and authorization parameters to enforce governance
- Online governance and instructions for data access (when data available)

 Cloud-based data available through governed APIs Data, metadata, code, tools and infrastructure available through public, versioned, open-source code on GitHub

Connect Data Ecosystem provides an interoperable infrastructure



Example of Front-End User Interface: MyConnect Participant App



Benefit of Middle Layer of APIs: Pulling from multiple resources

Baseline Recruitment

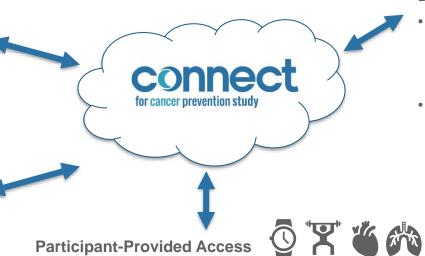


- Deidentified data on eligible members/patients
- Verification of consented individuals in MyConnect
- Study activity flags

Study Services



- Communications, Twilio
- Incentives
- Quest survey tool



Participant-Provided Access

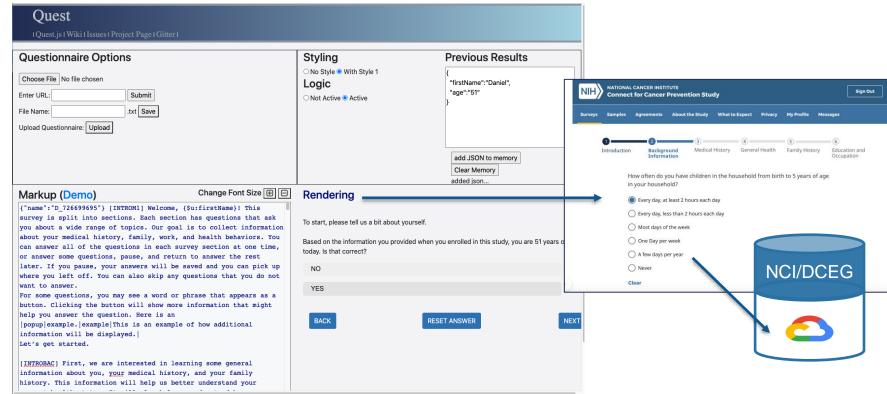
- Wearable devices: FitBit, air monitors, etc.
- External health care records: Medicare, other EHR for participants that leave partner healthcare systems, etc.

Data Linkages



- Health outcomes: cancer registries, virtual tumor registry, NDI mortality, HIV registry
- Geospatial data: outdoor air quality data, other EPA data, US Census

Quest provides open-source code to render surveys into MyConnect and capture data into Connect's backend data



Governance & Organizational Structure



Connect Governance

Connect Leadership and Coordinating Center

Governing Board

Stephen Chanock (Chair)

Executive Leadership

Steering Committee

Chief Scientist (Chair), DCEG Connect PIs, Coordinating Center Leadership, DCEG Associate Directors

Executive Committee

Chief Scientist (co-Chair), Rotating Site PI (co-Chair), Steering Committee Members, Site PIs, Participant Advisory Board Lead, Support Service Contractor Leads

External Advisory and Review Groups

International Scientific Advisory Board

Naomi Allen, Eric Boerwinkle, Julie Buring, Carmen Guerra, Chris Haiman, David Hunter, Roger Milne, Alpa Patel, Karriem Watson

National Cancer Institute (NCI) Board of Scientific Counselors (BSC) – Clinical Sciences and Epidemiology

Participant Advisory Board (PAB)

Connect Leadership and Organization

Governing Board

Executive Leadership

Steering Committee

Executive Committee

Coordinating Center

Mia Gaudet, Chief Scientist of Connect

Resource Access Committee

Representatives from the Connect Coordinating Center, Executive Committee, external investigators, study participants, ethics experts, and funding agencies

Ad hoc Expert Review Panels

Subcommittees

Biospecimens • Questionnaires • Devices Meta-Level Data • Medical Records

Ad hoc Working Groups

EpiSphere Data Science & Engineering Research Group

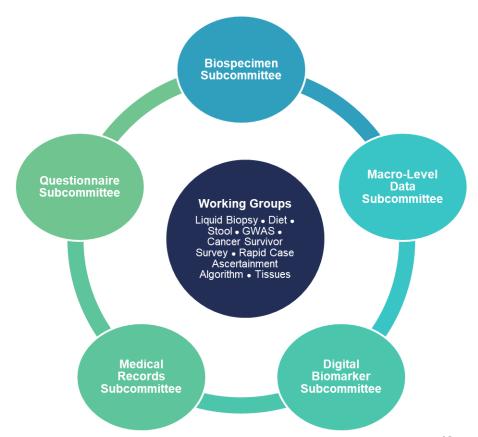
Jonas Almeida, Chief Data Scientist



Connect Resource Development Structure

Subcommittees (standing)

- Guide development of Connect infrastructure and enhancements
- Led by DCEG with NIH investigators
- Report to Steering Committee
- Working Groups (ad hoc)
 - Led by nominator with investigators from NIH, recruitment sites, and scientific community
 - Develop scientific and technical details of proposals and protocols
 - Report to relevant Subcommittees





Resource Enhancements

New questionnaire modules, biospecimens, other sources of data

Connect Resource Building Principles



Research resource enhanced by scientific community



Balance scientific value with participant burden

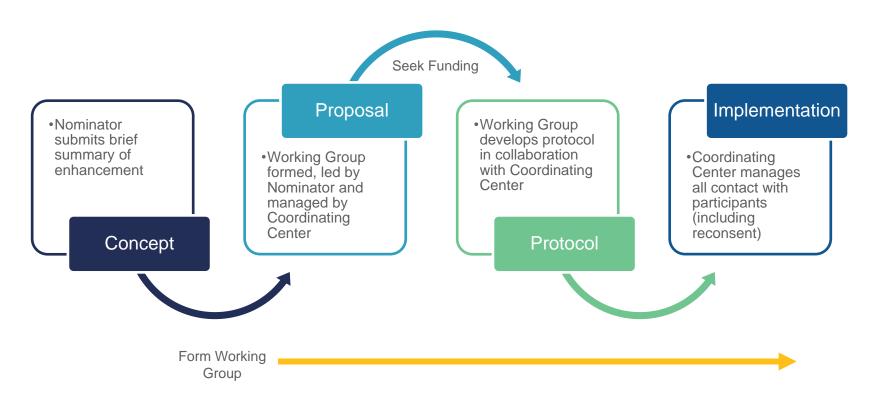


Maximize participant privacy and confidentiality



All enhancements grow Connect resources

Connect Resource Enhancement Approval Process





Resource Access Principles



Connect Resource Access Principles



Research resource for scientific community



Broad data sharing policies

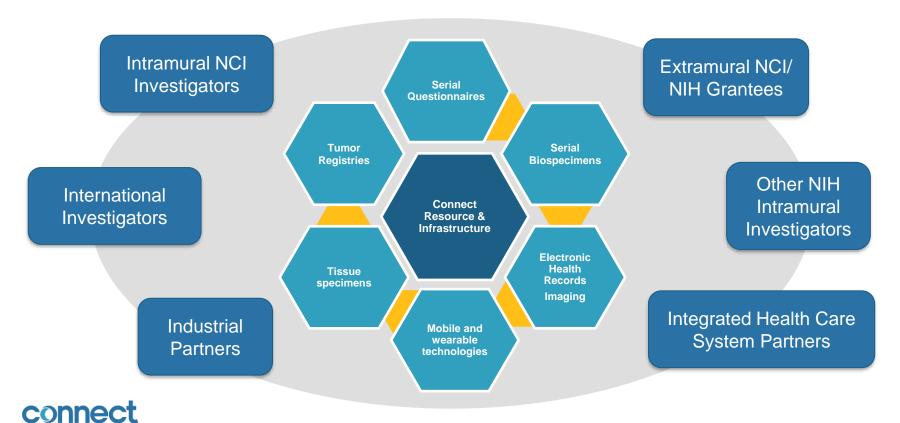


Participant privacy and confidentiality



F.A.I.R. data infrastructure

Resource Access for All*



Levels of Data Access

