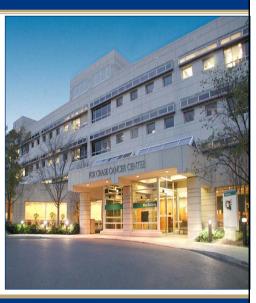
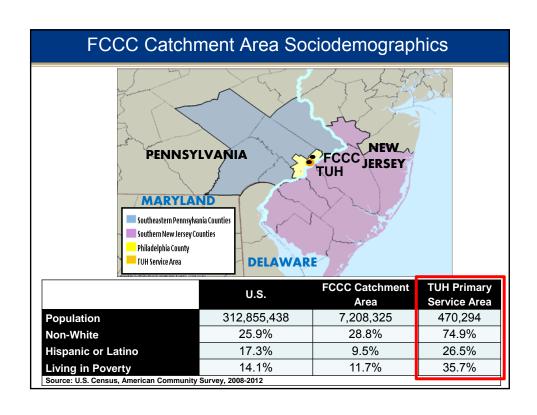
#### FCCC CCSG Supplement: Population Health Assessment

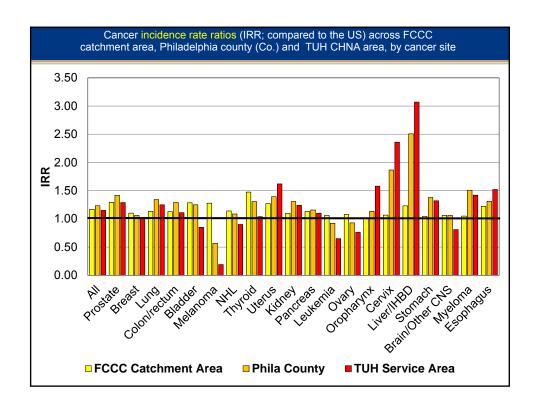
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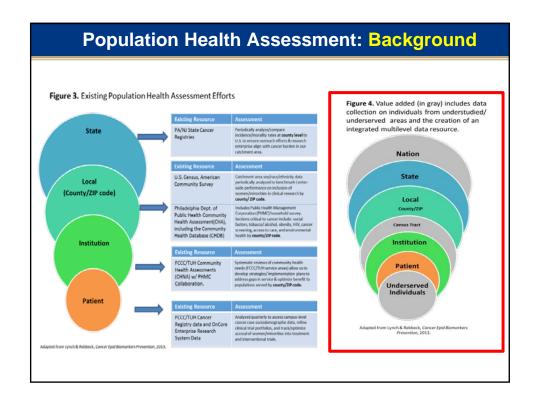






# Population Health Assessment: Background

- Capitalizing on existing population health assessment efforts/infrastructure, we propose a local data collection project that will allow for
  - · improved characterization of our catchment area
  - new insights into underserved/understudied populations
  - development of a novel, multilevel data resource relevant to national/state/local cancer surveillance efforts



# **Population Health Assessment: Study Population**

- Individually survey ≥ 1000 persons residing in our catchment area
  - ≥ 50% of the individuals surveyed will reside in the TUH Primary Service
     Area
  - ≥ 50% of the individuals surveyed *across the catchment area* will satisfy at least one of the following criteria of being historically underserved
    - 1) Live in medically underserved areas (Health Resources and Services Administration)
    - 2) Have low literacy
    - 3) Have low income (≤ \$11,670 for single individual or \$23,850 for family of 4
    - 4) Be uninsured/subsidized health insurance coverage

# Population Health Assessment: Recruitment Plan

- 1) We will identify/recruit subjects who reside in the TUH Primary Service Area by expanding Temple Health Block-by-Block's (THB3) already-planned, third wave of data collection
  - Re-contacting families already participating in THB3
- 2) We will identify/recruit (historically underserved and "nonunderserved") subjects across the entire catchment area (both within and outside the TUH Primary Service Area)
  - Via future community outreach/cancer screening efforts scheduled/conducted by FCCC's Office of Community Outreach (OCO)
  - Through the OCO's and the Health Communication and Community Engagement (HCCE) Facility's community partners
  - Re-contacting individuals who previously participated in research studies conducted by the HCCE Facility

## **Temple Health: Block-by-Block Project Goals**

...establish a multigenerational urban neighborhood health partnership to enrich minority research participation, optimize the institutional research portfolio, and contribute to the health improvement of our community.

- Provide a sustainable cohort of community participants engaged in individual and community health improvement across the life cycle through research;
- Provide investigators access to descriptive health data from urban populations for research purposes;
- Enhance participation of minorities in clinical research through improved education, access, and empowerment;
- Provide a venue for a community voice in new research initiatives.

## **Temple Health: Block-by-Block Recruitment Goals**

- 1,000 subjects per year
- Ages ≥ 18 years (children to be included in Year 3)
- · Multiple individuals within each household
- Maintain retention >80%
- Minimum goal of 5,000 subjects accrued and retained



## **Temple Health: Block-by-Block Methods**

- Collect descriptive, longitudinal data every 6 months related to:
  - Demographic and social characteristics
  - Anthropometric & biologic parameters
  - Health behaviors
  - Health outcomes
- Specific data "themes" for each bi-annual assessment
- Integration of data collection/interventions for independent research studies



## **Specific Benchmarks of Success**

- Increased minority recruitment into TUHS clinical trials
- Increased clinical investigations which focus on community health priorities
- Increased representation of minorities in national, therapeutic studies
- Improved health indicators (very long term ©) among North Philadelphia residents:
  - Jobesity and improved nutrition
  - Itobacco use
  - health screening
  - · earlier disease diagnosis



# **Population Health Assessment: Recruitment Plan**

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#### **Population Health Assessment: Key Constructs** Table 1. Data and key constructs for proposed data collection project. Data/Key Construct Instrument Instrument U.S. Census [1]\*, HINTS [2]\*, NHIS CCS [3]\*, BRFSS [4]\* Sociodemographics (e.g., age, gender, marital status, race, ethnicity, preferred language, education, income, urban/rural status, insurance status, etc) Reactions to race Social context (e.g., work, finances, etc) HINTS [2]\* BRFSS [4]\*, HINTS [2]\* Social networks Chronic health conditions/ functional status HINTS [2]\*, BRFSS [4]\* BRFSS [4]\*, SF-12 [5] HINTS [2]\* Health care access Health status/HRQOL Cancer perceptions/knowledge HINTS [2]" HINTS [2]", NHIS CCS [3]", BRFSS [4]" HINTS [2]", NHIS CCS [3]", BRFSS [4]" HINTS [2]", NHIS CCS [3]", BRFSS [4]" HINTS [2]", NHIS CCS [3]", BRFSS [4]", 13-item Attitudes Toward Genetic Research HINTS [2]\* Cancer communication Cancer-related risk behaviors Cancer risk attitudes Cancer screening Family history/genetic testing Knowledge/attitudes regarding medical research/clinical trials of Participation Scale [7] Knowledge/attitudes regarding biobanking research biobanking research biobanking research , denotes selected items; HINTS, Health Information National Trends Survey; NHIS CCS, National Health Interview Survey Cancer Control Supplement; BRFSS, Behavioral Risk Factor Surveillance System; SF-12, 12-Item Short Form Health Survey; BANKS, Biobanking Attitudes and Knowledge Survey

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