

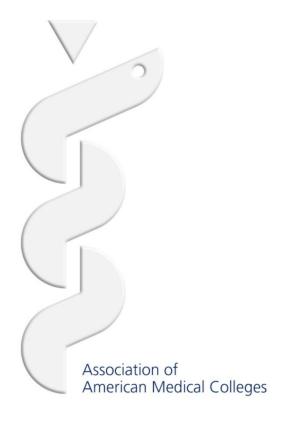


Promising Practices to Improve Hispanic Health

Collaborative interprofessional forum to discuss advancement of Hispanic health

Approaches to Hispanic Health Research

September 25, 2018 1:30-3:00 p.m. ET Learn Serve Lead



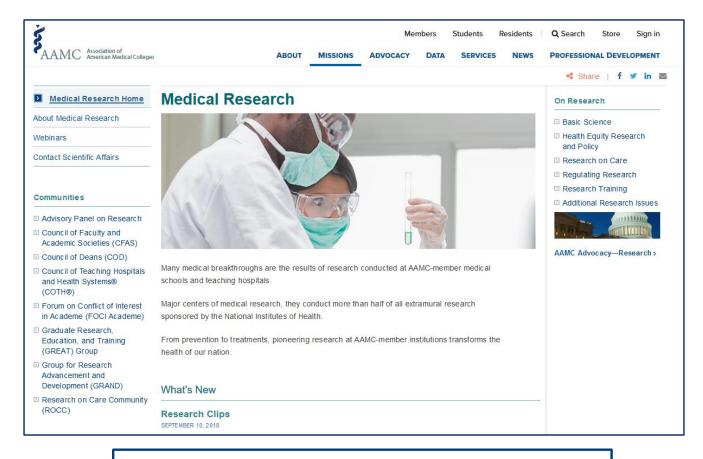
Welcome & Introductions: Malika Fair, MD, MPH



Senior Director, Health Equity Partnerships & Programs Association of American Medical Colleges



AAMC Resources: Website









AAMC Resources: Newsletters



Conferences, Meetings, and Webinars

Just Announced Happening in May and June

Grant Funding Opportunities

Just Announced

LOIs and Applications Due in May

Fellowship and Training Opportunities

Just Announced

LOIs and Applications Due in May

Resources

New Resources

Conferences, Meetings, and Webinars

Just Announced

The Health Information Environment as a Social Determinant of Health: Implications for Health Disparities and Opportunities for Intervention

July 18, 2019

Sacramento, CA

North American Housing & HIV/AIDS Research Summit IX Structural Interventions and **Ending the Epidemic**

August 1-3, 2018

Washington, DC



The AAMC Population Health Connect newsletter is a product of the AAMC Public Health Initiatives portfolio to increase awareness about population health activities and curricular resources relevant to the academic medicine community. To learn more about the AAMC's Public Health Initiatives, visit aamc.org/publichealth.

Upcoming Webinars & Conferences

May 2018

The Wrong Complexion for Protection: Race, Place and the Politics of Pollution May 4, 2018

College Park, MD

3rd Annual NIH Vivian W. Pinn Symposium - Leveraging the Network to Advance Women in Science

May 16, 2018 Bethesda, MD

11th Annual National Conference on Health Disparities

May 16-19, 2018 Philadelphia, PA

2018 Wisconsin Public Health Association-Wisconsin Association of Local Health Departments and Boards Public Health Conference; Leading the Evolution of Public Health

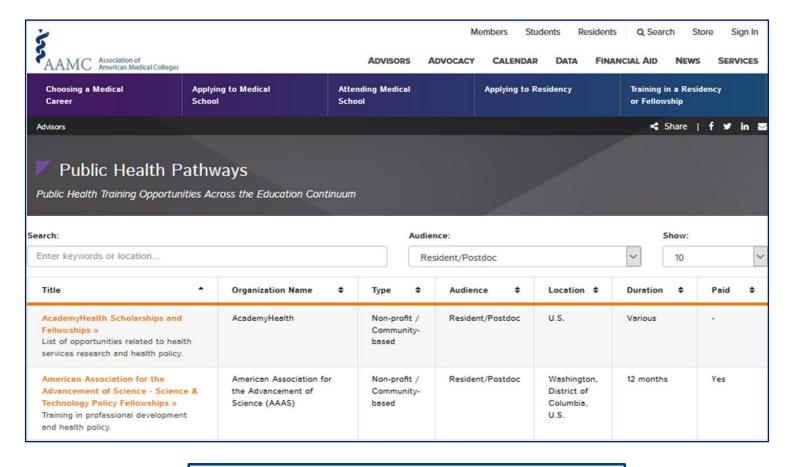
May 22-24, 2018 Green Bay, WI

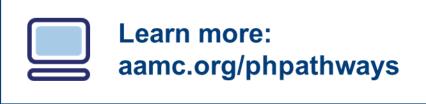
The Group on Diversity & Inclusion Professional Development Conference and 10th Anniversary Celebration - Tomorrow's Leaders: Bridging Boundaries Toward Equity





AAMC Resources: Public Health Pathways









Moderator: Maureen Lichtveld, MD, MPH Tulane University School of Public Health and Tropical Medicine



- Chair, department of global environmental health sciences
- Freeport McMoran Endowed Chair in environmental policy
- Director, Center for Gulf Coast Environmental Health Research, Leadership, and Strategic Initiatives















HISPANIC-SERVING HEALTH PROFESSIONS SCHOOLS

The Hispanic-serving Health Professions Schools (HSHPS) is composed of academic institutions that strive to achieve equitable health and well-being for our Hispanic community.



HSHPS RESOURCES

Graduate Fellowship Training Program

GFTP provides training opportunities for students and recent graduates interested in working on Hispanic health research. Fellows are placed throughout the United States and Latin America within government agencies and academic institutions. All HSHPS fellows work alongside a mentor, assist with a research project as it relates to minority health issues, and participate in a lecture series. Applications for Summer 2019 will open in November!

Networking

Our network not only consists of HSHPS members, but also federal agencies and non-profit organizations. Through our network, Member Institution's have an opportunity to collaborate on research at a local or national level, provide or seek mentorship, and more.

e-Newsletter

Our Member Institutions receive a quarterly newsletter which includes: HSHPS activities, legislative updates, job and funding opportunities, upcoming events, and spotlight a member faculty and student who have contributed to the field of Hispanic health or Hispanic health workforce through scholarly research.

Webinars

Member Institutions receive complimentary access to attend quarterly HSHPS educational webinars. Webinars are designed to create a collaborative inter-professional forum to discuss advancement of Hispanic health through community engagement, academic medicine, medical education and clinical approaches to address health disparities.











Learning Objectives

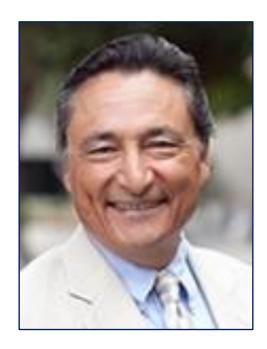
- Discuss current practice approach to Hispanic research at the community, national and global levels
- Describe intervention trends in reducing Hispanic health disparities
- Examine common challenges and barriers in Hispanic health research



Panelists



Norma A. Pérez, MD, DrPH
President, HSHPS
Executive Director, Hispanic Center of
Excellence
University of Texas Medical Branch



Gregory A. Talavera, MD, MPH
Professor
South Bay Latino Research Center
San Diego State University School of
Public Health



Pierre Buekens, MD, MPH, PhD
W. H. Watkins Professor
Director, CERPE
Tulane University School of Public Health and
Tropical Medicine





Norma Alicia Pérez, MD, DrPH University of Texas Medical Branch



- Executive Director, Hispanic Center of Excellence (HCOE) and Special Programs
- Faculty appointments with Internal Medicine-Geriatrics and Preventive Medicine & Community Health
- President, Hispanic-Serving Health Professions Schools



Promising Practices to Improve Hispanic Health

Collaborative interprofessional forum to discuss advancement of Hispanic health

Community research partnerships: Are we true partners?

Norma A. Pérez, MD, DrPH
President, Hispanic Serving Health Professions Schools
Director, School of Medicine Special Programs and Hispanic Center of Excellence
Assistant Professor of Internal Medicine-Geriatrics and Preventive Medicine and Community Health



Working together...

No matter how much success you're having, you can't continue working together if you can't communicate.

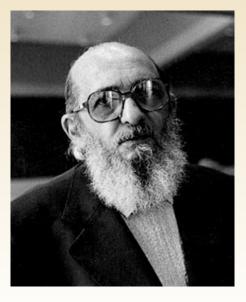
- Matt Cameron



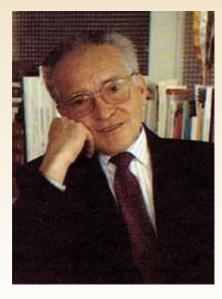
Pioneers



Kurt Lewin



Paulo Freire



Orlando Fals

Strong Community-based Participatory Research Funders

2001 - Institute of Medicine

Centers for Disease Control

National Institutes of Health

National Institute of Environmental Health Sciences

Agency for Healthcare Research and Quality



Three Models of Community-based Participatory Research

Model 1

Single-theme collaboration between a university-based center and a community organization

Model 2

Targeted, areabased collaboration between academic researchers and a small group of community organizations

Model 3

Broad-based coalition of grassroots organizations



Ethical Challenges

The Principle of Beneficence

Kelly's Principals

Camara Jones' typology



COMMUNITY TOOL BOX

It is a public service of the Center for Community Health and Development at the University of Kansas. It is a free, online resource for those working to build healthier communities and bring about social change.

Since 1994

Over 300 educational modules and other free tools

Available in English, Spanish, Arabic, and Farsi, and with millions of user sessions annually, it has reached those working in over 230 countries around the world.

https://ctb.ku.edu/en/about-the-tool-box



Local Best Practice of Community-based Participatory Research Program





The ITS, established in 2007, is the academic home of UTMB's Clinical Translational Sciences Award (CTSA), funded by the National Institutes of Health's National Center for Advancing Translational Sciences (NCATS)

Roles for Community Engagement

Engage communities around specific projects and advance the science of community engaged research

Identify community health needs and priorities

Provide input on research questions relevant to the community

Contribute to appropriate research design and methods

Develop culturally sensitive and ethical proposals

Enhance the recruitment and retention of research participants

Implement and disseminate research findings effectively

Provide culturally competent training for staff and communities

Foster innovation and participate in the dissemination of best practices to other CTSA Program Hubs



What Does Success Look Like?

Communities participate fully across the translational spectrum

Communities are consulted for their research priorities

Stakeholders are respected, valued, and rewarded

Effective implementation and dissemination strategies are developed

Increased public support for research

Improvements in the health and well-being of communities

Community engagement integrated into leadership, research, and communication strategies

Community engagement addressed as a scientific problem to identify best practices and most effective approaches

Community engaged research is rewarded in academic policies



Thank you



Gregory A. Talavera, MD, MPH San Diego State University Graduate School of Public Health



- Professor of Public Health, Division of Health Promotion & Behavioral Sciences
- Co-director, South Bay Latino Research Center
- Previously practiced ambulatory medicine in the Spanish-speaking, underserved communities of San Diego's border region





Latino Longitudinal Cohorts
-All of Us' Research Program
-Hispanic Community Health Study



Gregory Talavera, MD, MPH, Professor Health Promotion and Behavioral Science, Graduate School
of Public Health, San Diego State University



The Hispanic Community Health Study – Study of Latinos Overview, Baseline Results & Potential Disparities Implications

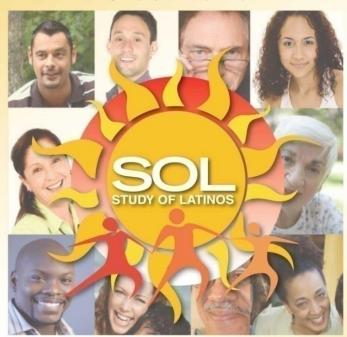
Gregory A. Talavera, MD, MPH
Professor San Diego State University
PI for the San Diego Field Site

ESTUDIO DE LOS LATINOS

Hispanic Community Health Study
STUDY OF LATINOS

¡Ayúdenos a entender la salud de nuestra gente!

Si tiene de 18 a 74 años de edad y es hispano/latino, usted puede ser parte de este histórico estudio.



Help us understand the health of our community!

If you are Hispanic/Latino between 18-74 years old, you can be part of this historical stud-

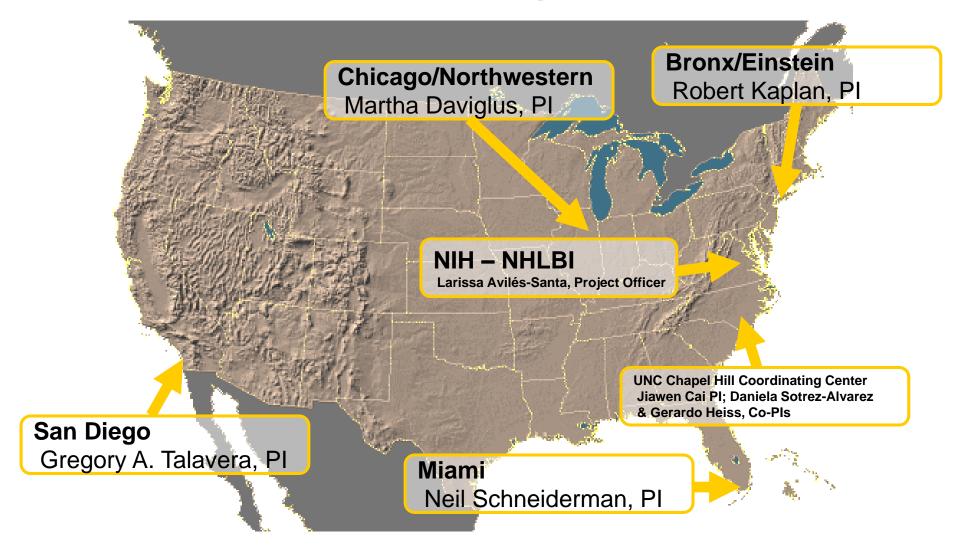
Primary Goals



- To identify the prevalence of cardiovascular and pulmonary disease and other conditions in U.S. Hispanic/Latino groups of <u>diverse</u> <u>backgrounds</u>.
- To identify the **prevalence of factors** that protect from or increase the risk for cardiovascular and pulmonary disease and other conditions in diverse U.S. Hispanic/Latino groups.
- To identify all-cause mortality, and the incidence of fatal and non-fatal cardiovascular and pulmonary events in diverse U.S. Hispanic/Latino groups. (2014) Birth outcomes



HCHS/SOL Field Centers & Coordinating Center



16,418 Participants

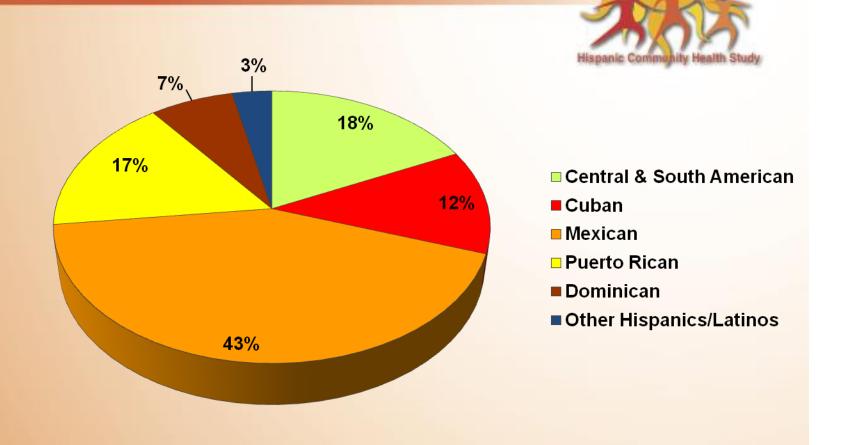
Random sampling of the local population Representative of the local communities

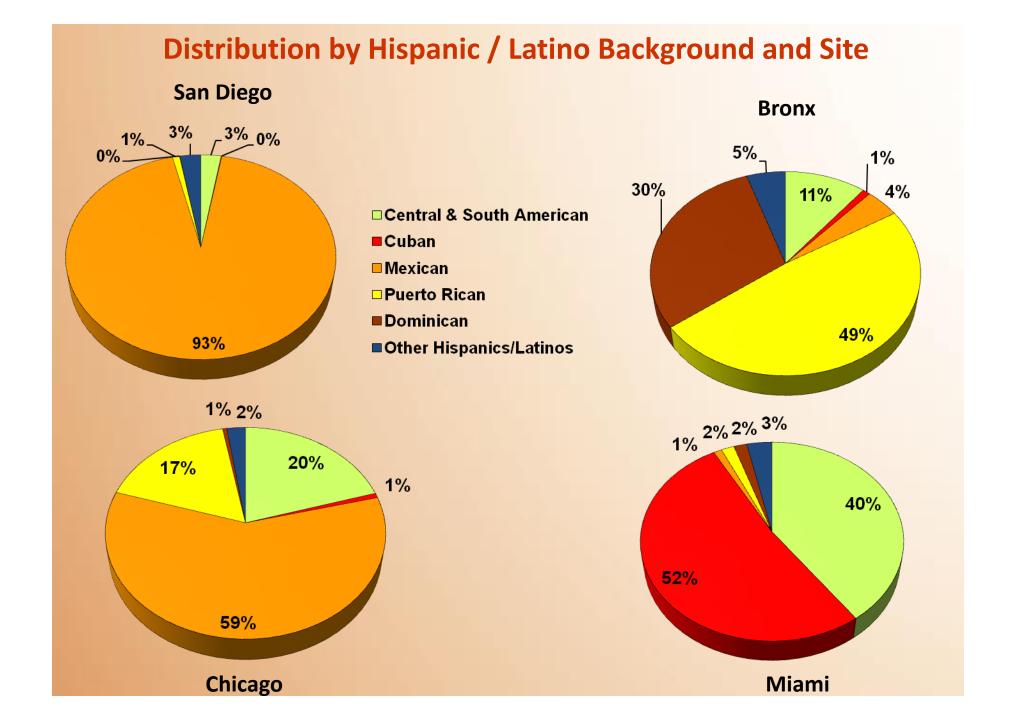
- Ages: 18 -74 years
 - **6,000** ages 18-44 years
 - **10,000** ages 45-74 years
- Approximately 4,000 persons per Field Center who self-identify with any of the following Hispanic/Latino heritage/background groups:
 - Mexican background
 - Puerto Rican background
 - Cuban background
 - Dominican background
 - Central/South American background
 - Other Hispanics/Latino background groups

Other Longitudinal Studies



Distribution of Study Cohort at Baseline by Hispanic/Latino Heritage/Background

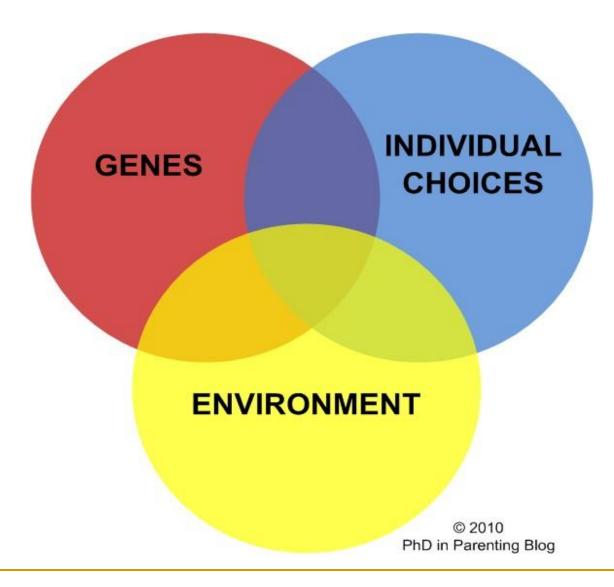




Publications Parent Study

~500 Manuscripts
487 manuscripts in concept phase

Gene-Environment-Behaviors



Continental Ancestry

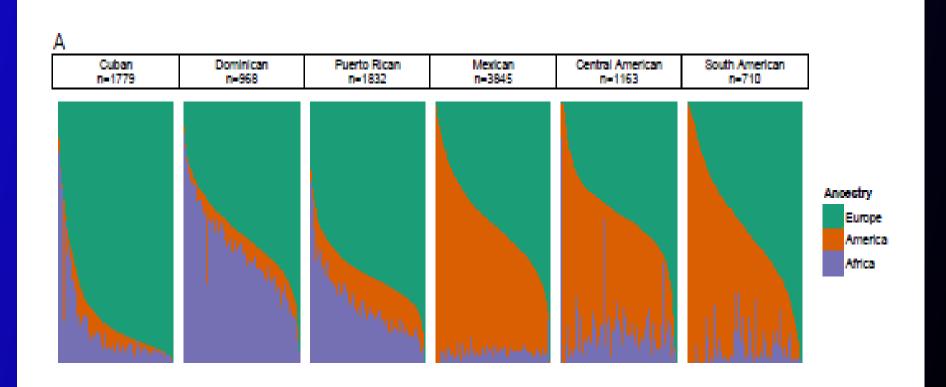


Figure 2: Continental ancestry proportions for the autosomes and X chromosome

(A) Estimates of continental ancestry proportions on the autosomes for an unrelated set of HCHS/SOL individuals grouped by self-identified background, with the number of unrelated individuals shown for each group. Each vertical bar represents a single individual, within which the three ancestry fractions are represented by three color-coded segments.

Prevalence of CVD Risk Factors by Hispanic/Latino Group - MEN

CVD Risk Factors (%) ^a		Hispanic/Latino Group						
	All	Cuban	Dominican	Mexican	Puerto Rican	Central Am.	South Am.	
Unweighted N	<i>5979</i>	1034	480	2337	1067	648	413	
Hypertension	25.4	28.9	32.6	21.4	27.4	25.0	19.9	
Hypercholesterolemia	51.7	53.7	47.6	53.9	48.2	54.9	52.2	
Obesity	36.5	33.6	38.6	36.8	40.9	32.7	26.8	
Diabetes	16.7	13.2	18.2	19.3	16.2	16.3	10.1	
Smoking	25.7	31.1	11.1	23.1	34.7	19.9	15.1	

^a Values (except N) weighted for survey design and non-response ,and age standardized to Census 2010 US population. **Hypertension** SBP/ DBP >140/>90 or on treatment. **Hypercholesterolemia**, total cholesterol >240 mg/dL HDL-C <40 mg/dL LDL-C >160 mg/dL or on treatment. **Obesity,** BMI >30kg/m2; **Diabetes**, fasting glucose >126 mg/dL 2h-post-load plasma glucose >200 mg/dL A1c >6.5%, or use of diabetes medications. **Smoking**, currently smoking cigarettes.

Prevalence of CVD Risk Factors by Hispanic/Latino Group - WOMEN

CVD Risk Factors (%) ^a		Hispanic/Latino Group						
	All	Cuban	Dominican	Mexican	Puerto Rican	Central Am.	South Am.	
Unweighted N	9100	1167	920	3895	1523	986	609	
Hypertension	23.5	26.4	26.1	19.5	29.1	25.6	15.9	
Hypercholesterolemia	36.9	37.5	33.1	36.2	41.0	39.4	31.4	
Obesity	42.6	38.9	42.5	41.5	51.4	41.6	30.8	
Diabetes	17.2	13.9	18.0	18.5	19.4	17.9	9.8	
Smoking	15.2	21.2	11.7	10.0	31.7	8.7	11.3	

aValues (except N) weighted for survey design and non-response and age standardized to Census 2010 US population. **Hypertension** SBP/ DBP >140/>90 or on treatment. **Hypercholesterolemia**, total cholesterol >240 mg/dL HDL-C <40 mg/dL LDL-C >160 mg/dL or on treatment. **Obesity,** BMI >30kg/m2; **Diabetes**, fasting glucose >126 mg/dL 2h-post-load plasma glucose >200 mg/dL A1c >6.5%, or use of diabetes medications. **Smoking**, currently smoking cigarettes.

Overview of the *All of Us* Research Program





Gregory Talavera, MD, MPH, Professor Health Promotion and Behavioral Science, Graduate School of Public Health, San Diego State University

What is precision medicine?

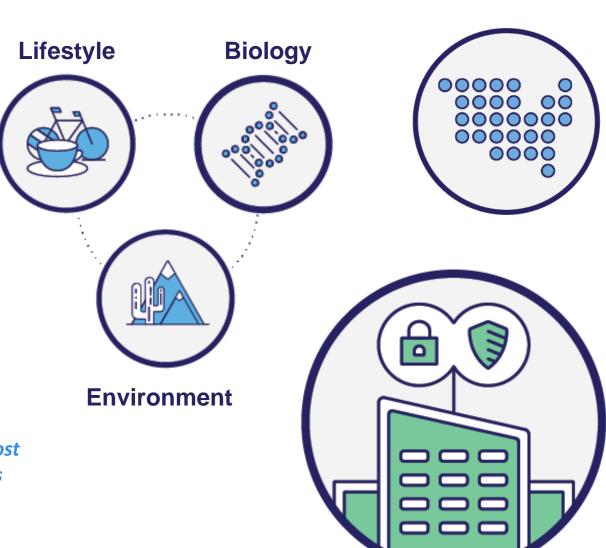
Precision medicine is an emerging approach for disease treatment and prevention that takes into account individual variability in lifestyle, socioeconomics, environment, and biology.

It is a radical shift in how each of us can receive the best care possible based on our unique makeup.



"All of Us is among the most ambitious research efforts that our nation has undertaken!"

NIH Director Francis Collins, M.D., Ph.D.



Why do we need the All of Us Research Program?



People/ Patients

- Patients may not be served well by treatments designed for the "average" patient.
- Health problems can take years to unravel and require much trial and error treatment.
- Patients may not have access to or make use of their own health data.



Professional Providers

- Precision medicine is still in the early days, so providers do not have enough information available to provide precision care for many conditions.
- Developing individualized approaches to care often requires overtime.
- It can be difficult to coordinate care between many different providers, especially with medical records and key data scattered in different silos.



Biomedical Researchers

- Researchers spend a lot of time and resources creating new IT systems, databases, and analytic tools.
- Data collection is not standardized so data can be siloed and difficult to integrate.
- A single lab's resources may not be sufficient to answer the research question.

National Network of Inaugural Partners

This map is a visual representation of the large and nationally distributed community of partners to the *All of Us* Research Project.





Kinds of Questions this Resource May Help Answer

How can we prevent the chronic pain that affects more than 100 million people across the U.S. each year?

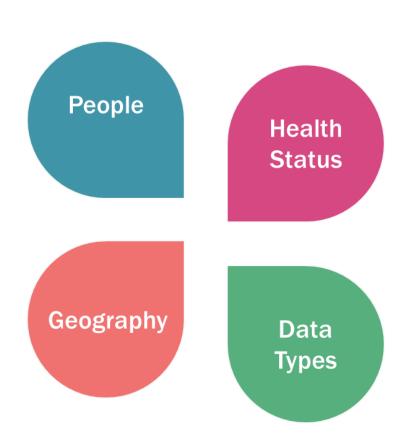
Or develop better pain medicines that aren't addictive?

Or develop better treatments for diabetes, which affects almost 10% of Americans—or prevent diabetes altogether? Or slow or even stop different kinds of dementia?

Or develop more cancer cures that will work the first time, so we can skip painful trial-and-error chemotherapy?

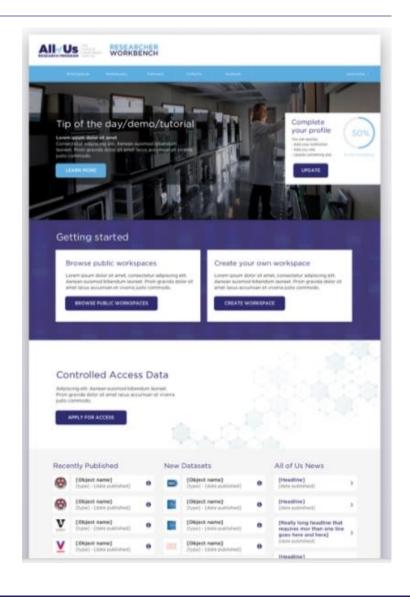
Innovative Aspects of All of Us

- Diversity at the scale of 1 million people: demographically, geographically, medically, and especially those underrepresented in biomedical research
- Diversity of data types collected longitudinally: clinical, environmental, genetic, behavioral, socioeconomic
- Focus on participants as partners: included in governance, invited to co-invent systems and give input into the science, choice to receive all data and information back
- National, open resource for all: open to the public and all researchers, open source software & tools



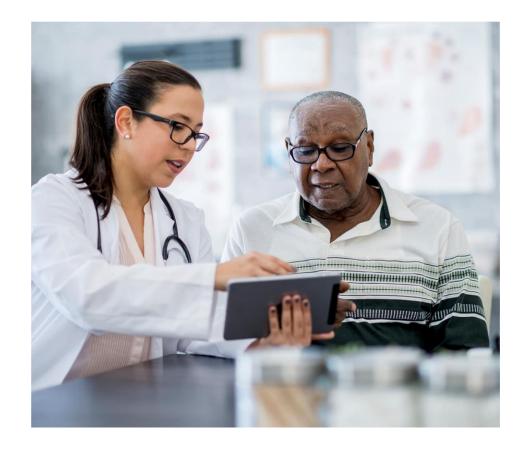
What is the promise for researchers?

- The opportunity to save time and resources and accelerate your research breakthroughs by leveraging:
 - A **rich resource of data**, including biospecimens and increasingly robust electronic health records.
 - A longitudinal dataset that will follow participants as they move, age, develop relationships, get sick, and try treatments.
 - A diverse cohort of participants, including people both healthy and sick, from all walks of life and all parts of the country.
 - Both raw data and data that is already cleaned and curated.
 - Robust computing and analytic tools to support complex data analyses in a secure data environment.
 - A group of engaged participants who may be eager to participate in ancillary studies.
- The ability to easily share workspaces and analyses with research partners and reviewers.
- The chance to learn from the program's pilots and experiments and leverage innovations for other studies and cohorts.



What is the promise for providers?

- Over time, increased scientific evidence and improved guidelines to enable precision medicine opportunities for more people and conditions:
 - Better understanding of the impact of environment and lifestyle factors on health.
 - Increased knowledge of differences in risk factors and response to treatments among diverse populations.
 - More information on the development of conditions that will allow for earlier detection.
 - Deeper understanding of different conditions that may allow for **better stratification**.
- Innovations that may make it easier to share electronic health records with other providers and patients.
- New knowledge to help address health disparities, increase patient engagement, and understand the usefulness of consumer health devices and apps.



Two Methods of Enrollment



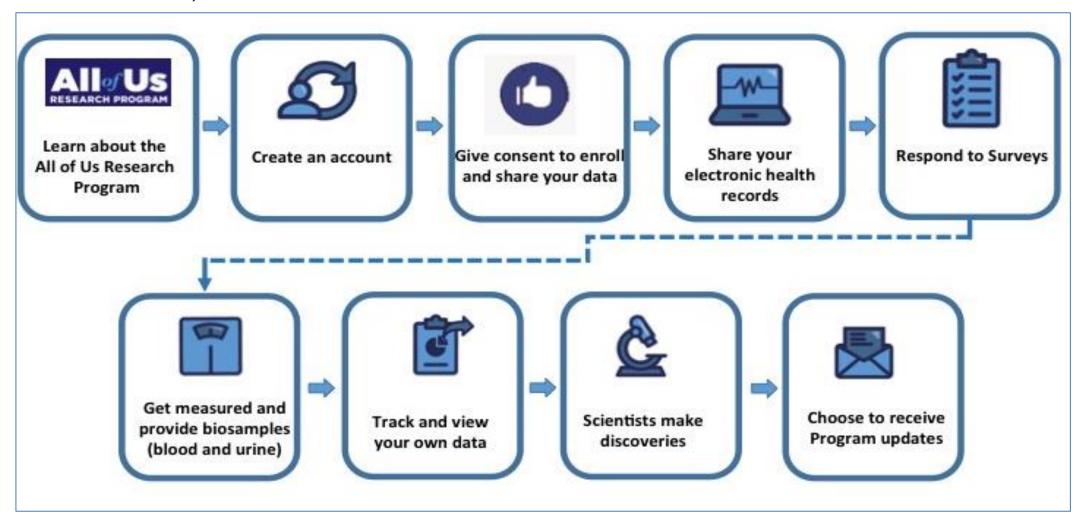
DIRECT VOLUNTEERS



HEALTH CARE PROVIDER ORGANIZATIONS

The Participant Journey

(Figure 6.1 of Core Protocol V1.7)





All of Us Research Program Data

The program will start by collecting a limited set of standardized data from sources that will include:

- Participant surveys
- Electronic health records
- Physical measurements
- Biosamples (blood and urine samples)
- Mobile/wearable technologies
- Geospatial/environmental data

Data types will grow and evolve with science, technology, and trust.



Minorities make up

380/0
of the US population.

Minority populations to rise to over

56% of overall population.

Minority enrollment in clinical trials?

>10%

VALUE TO PARTICIPANTS

African American population = 13.2% of US population

5% of clinical trial participants nationwide

Hispanic population = 17% of US population

- 7.6% of NIH clinical trial participants
- 1% of clinical trial participants nationwide

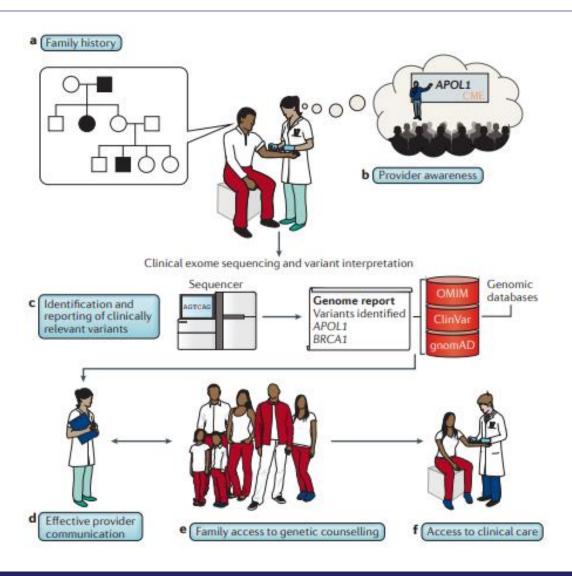
Underrepresented in biomedical research populations include ethnic minorities, as well as age, sex, gender, orientation, income, education, geography, access to care and disability.

A Transformational Approach to Participation

Participants in the *All of Us* Research Program will be true partners—not patients, not subjects—in the research process. Involved in every step of program development:

- What data we collect
- What lab analyses we do
- What research is conducted
- How data gets returned

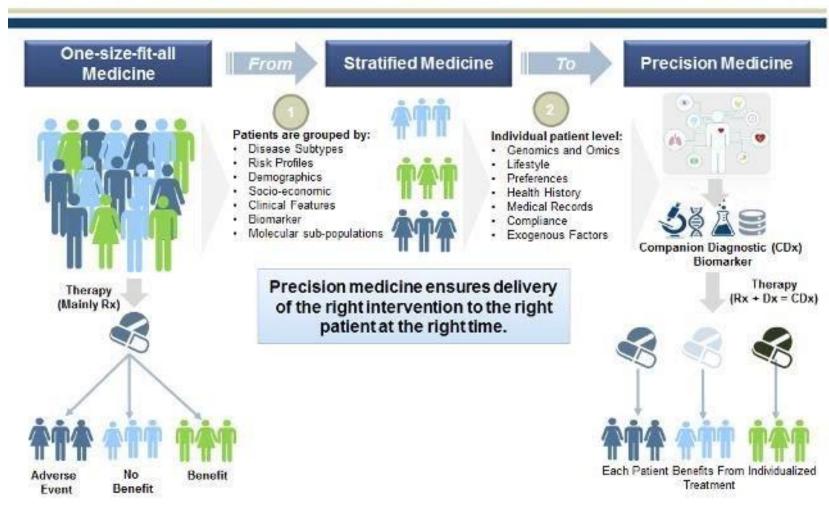




Pharmacogenomics

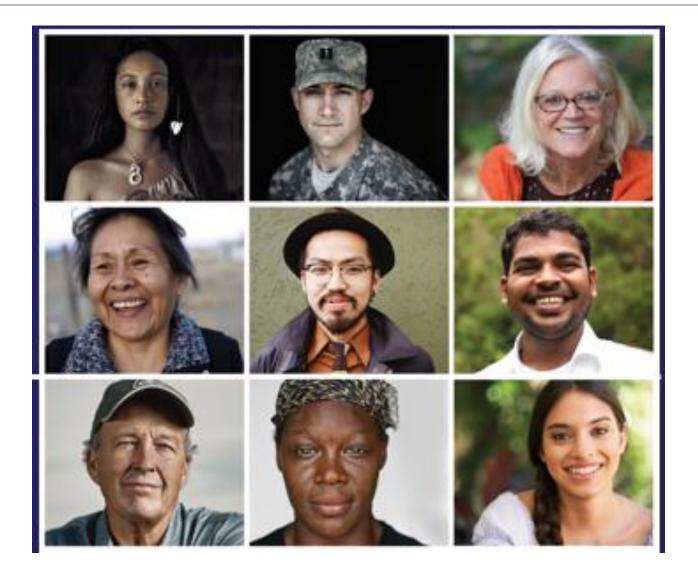
New Paradigm Shift in Treatment

Transitioning From the 'one-size-fits-all' to 'precision medicine' model with multi-level patient stratification.



Drug Industry Bets Big On Precision Medicine: Five Trends Shaping Care Delivery

Thank You



Pierre Buekens, MD, PhD Tulane University School of Public Health and Tropical Medicine



- W. H. Watkins Professor in the Department of Epidemiology
- Director, Tulane Center for Emerging Reproductive and Perinatal Epidemiology (CERPE)
- Co-founded Trop-G, a collaboration working to control and eliminate NTDs in high-risk areas around the Gulf of Mexico





Accessing Mexican Literature and Data

Pierre Buekens, MD, PhD





Accessing Mexican Literature

- Virtual Library: http://bvsalud.org
- Lilacs: http://lilacs.bvsalud.org
- Scielo: http://www.scielo.org.mx

bvsalud.org







Latin American and Caribbean Center on Health Sciences Information



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LILACS is the most important and comprehensive index of scientific and technical literature of Latin America and the Caribbean. For 32 years contributing to increase visibility, access and quality of health information in the Region.

LILACS in numbers

26 countries

918 journals

827.400 records

690.750 articles

87.713 monographies

40.647 thesis

418.391 full texts





Polling Question

What is the best Mexican Public Health Journal?

- A. BJX Salud Pública
- B. QRO Salud Pública
- C. Salud Pública de México
- D. There is no Mexican Public Health Journal

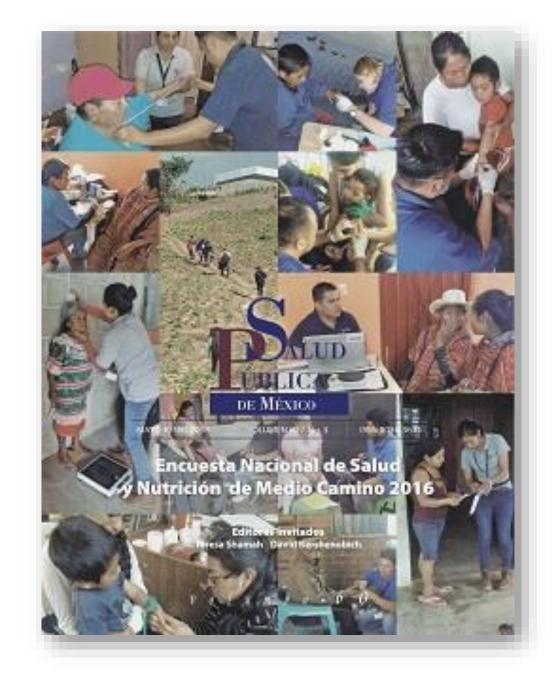


Salud Pública de México











Mexican Surveys and Databases

- https://www.gob.mx.conapo
- http://www.inegi.org.mx
- http://www.dgis.salud.gob.mx



Low Birthweight in Mexico: A Systematic Review

Pierre Buekens · Caitlin Canfield · Nicolas Padilla ·

Elia Lara Lona · Rafael Lozano

Surprisingly, most of the US literature does not refer to birth outcome data available in Mexico.



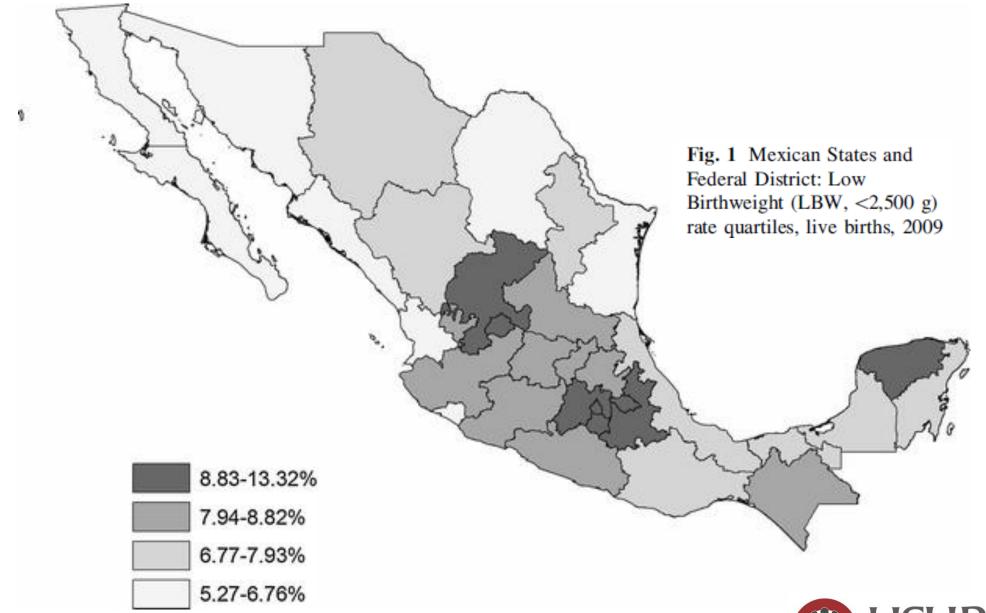


Table 2 continued

State and City (if applicable)	Sample type	Year	Sample size (# of births or women*)	LBW %
National-Level Survey (DHS/ENFES)	National Household Survey	1982–1987	5,311	10.2
National-Level Survey (ENADID)	National Household Survey	1992–1997	23,607	11.0
National-Level Survey (ENSAR)	National Household Survey	2003	19,612*	10.0



Conclusions

- Mexican literature should be cited
- Results from Mexican surveys should be used
- Mexican data available online should be analyzed in collaboration with Mexican colleagues.



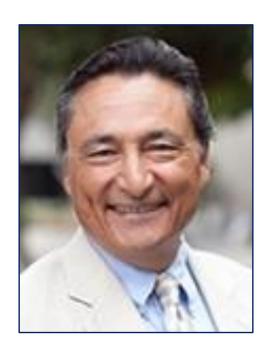
Center for Emerging Reproductive and Perinatal Epidemiology (CERPE)



Approaches to Hispanic Health Research: Q&A



Norma A. Pérez, MD, DrPH
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Tropical Medicine







Learn

Serve

Lead

Association of American Medical Colleges