

Massachusetts Department of Public Health

COVID-19 Community
Impact Survey (CCIS): Reflecting on
Impact & Looking Ahead

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Presented by Lauren Cardoso, PhD

CCIS TEAM MEMBERS

CCIS Project Leads

W.W. Sanouri Ursprung, Lauren Cardoso, Beth Beatriz, Glory Song, Caroline Stack, Kathleen Fitzsimmons, Emily Sparer-Fine, Ta-wei Lin, Lisa Potratz, Heather Nelson, Amy Flynn, Lisa Arsenault, Abby Atkins

CCIS Steering Committee

Lauren Cardoso, W.W. Sanouri Ursprung, Beth Beatriz, Abbie Averbach, Ruth Blodgett, Ben Wood, Sabrina Selk, Jessica del Rosario Nicole Daley, Lisa Potratz

CCIS Analytic Team, Data to Action Team, Data Dissemination Team, Communications Team

Allison Guarino, Andrea Mooney, Angela Laramie, Ann Marie Matteucci, Anna Agan, Arielle Coq, Barry Callis, Beatriz Pazos Vautin, Ben Wood, Brittany Brown, Chelsea Orefice, Dana Bernson, David Hu, Dawn Fukuda, Ekta Saksena, Elise Pechter, Emily White, Fareesa Hasan, Frank Gyan, Glennon Beresin, Hanna Shephard, Hannah Walters, Hermik Babkhanlou-Chase, James Laing, Jena Pennock, Jennica Allen, Jennifer Halstrom, Justine Egan, Kathleen Grattan, Kim Etingoff, Kirby Lecy, Lamar Polk, Lauren Fogarty, Lauren Larochele, Mahsa Yazdy, Marianne Mabida, Matthew Tumpney, Megan Hatch, Megan Young, Melody Kingsley, Michelle Reid, Miriam Scrivener, Nassira Nicola, Nicole Daniels, Nicole Roos, Rebecca Berger, Rebecca Han, Robert Leibowitz, Susan Manning, Thomas Brigham, Timothy St. Laurent, Vera Mouradian, Victoria Nielsen, Ziming Xuan, Elizabeth Showalter, Priyokti Rana, Mayowa Sanusi, Emily Lawson, Alana LeBrón

CCIS COMMUNITY PARTNERS

Many groups that were critical in the success of this effort and gave important input on the development and deployment of the survey:

- Academic Public Health Volunteer Corps and their work with local boards of health and on social media
- Mass in Motion programs, including Springfield, Malden, and Chelsea
- Cambodian Mutual Assistance
- The Mashpee Wampanoag Tribe
- The Immigrants' Assistance Center, Inc
- Families for Justice as Healing
- City of Lawrence Mayor's Health Task Force
- The 84 Coalitions, including the Lawrence/Methuen Coalition
- Boys and Girls Clubs, including those in Fitchburg and Leominster and the Metro South area
- Chinatown Neighborhood Association
- Father Bill's
- UTEC
- MassCOSH
- Stavros Center for Independent Living
- Greater Springfield Senior Services
- Center for Living and Working
- DEAF, Inc.
- Massachusetts Commission for the Deaf and Hard of Hearing
- Viability, Inc.

Why did we conduct the COVID-19
Community Impact Survey (CCIS)?

BACKGROUND

Context

The pandemic is exacerbating pre-existing public health concerns and creating new health crises to address. Even people who have not become sick with COVID-19 are managing stress, uncertainty, and isolation during this challenging time. DPH and its partners need real time data to prioritize resources and inform policy actions.



Goal

DPH conducted a survey to understand the specific needs of populations that have been disproportionately impacted by the pandemic, including its social and economic impacts.



Actions

DPH will use and share these data to prioritize our pandemic response and to create new, collaborative solutions with community partners.



How did the CCIS fill these data gaps?

PRECISION PUBLIC HEALTH DATA NEEDS



Timely & nimble enough
to capture emerging
health needs



Detailed enough to illuminate
not just disparate outcomes
but also actionable root
causes of these inequities



Granular enough to tell us
where to tailor efforts to
certain geographies and
populations



Cross cutting enough to inform
how related content areas
across the department
intersect and should be
coordinated

CCIS APPROACH

- Conducted a self-administered online survey (fall 2020) with over **33,000** adults and **3,000** youth in the final sample
 - Paired with population specific focus groups
- Covered a wide range of topics specific to adults and youth respectively
 - Perceptions & experiences of COVID-19, Basic needs, Access to healthcare, Pandemic-related changes in employment, Mental health, Substance use, and Safety
- Available in 11 languages; additional focus groups also conducted in ASL
- Open ended questions captured previously unknown needs and barriers
- Weighted results to the state average, with different weights applied to youth and adult samples
- Recruitment via network of community-based organizations (CBOs)
- Employed a snowballing sampling strategy to ensure we reach key populations
 - eg. People of color, LGBTQ+ individuals, People with disabilities, Essential workers, People experiencing housing instability, Older adults, and Individuals living in areas hardest hit by COVID-19

DATA INNOVATIONS



Community engagement at every point ensures better questions, answers, and interpretation (eg. question development, pilot testing, recruitment, focus groups, dissemination)



Developed a **novel weighting/sampling approach** scalable across DPH to generate **granular** results



Mixed methods – focus groups and open responses allowed us to hear more nuanced stories and unknown health needs.



Population focused not condition focused- high representation by race, ethnicity, sexual orientation, gender identity, transgender status, types of disability, education, language spoken, industry/occupation, geography, employment status, age, etc.

ACTION INNOVATIONS



Built in Action workstreams (eg. data to action workgroup; public data access workgroup; community data technical assistance; contextualized webinars; and web reports.



Build racial and social justice framing and call to action to aid external audiences in acting on findings



Built in community engagement infrastructure to rapidly get direct community input and get data back to community



Sheds light on light on both “who” and “why” MA residents have been impacted, not just the “what” – (captures intersection of multiple root causes like occupation; SDOH: childcare; ability to socially distance)

Did it work?

OUR EFFORTS WERE SUCCESSFUL

- Unprecedented sample sizes allow results by number of groups including:
 - race, ethnicity, substance use history, incarceration history, sexual orientation, gender identity, transgender status, types of disability, income, education, language spoken, industry/occupation, geography, employment status, age, etc.
- Compared to past surveillance surveys, CCIS priority population samples reached:
 - **10x** as many Alaska Native/Native Americans
 - **10x** as many LGBTQ respondents
 - **5x** as many residents who speak languages other than English
 - **5x** as many Hispanic residents
 - **5x** as many Asian residents
 - **Over twice** as many respondents in other populations including the deaf/hard of hearing and Black community

*example comparison rates were calculated in comparison to the 2019 Behavioral Risk Factor Surveillance Survey (BRFSS) sample sizes

SOME KEY POPULATION-FOCUSED FINDINGS



Black and American Indian / Alaska Native youth were 2x as likely to lose someone close to them due to COVID-19, compared to all youth



1 in 3 Parents faced housing insecurity (50% more than respondents not parents)



1 in 5 Cambodian respondents lost their job (2nd highest among all CCIS ethnic groups)



70% of non-binary adults reported experiencing poor mental health 15+ days in the past month and 84% of non-binary and queer youth reported feeling sad or hopeless for 2+ weeks in past year



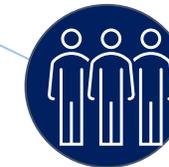
Respondents with a cognitive disability were 2x as likely to worry about getting food or groceries, compared to respondents without a cognitive disability



American Indians / Alaska Natives were 2.3x as likely to worry about getting access to broadband than White NH respondents



Hispanic/Latinx adults were 2x as likely as White Non-Hispanic adults to change the status or nature of their employment to take care of children.



1 in 4 Asian, Multiracial, and Black respondents experienced racial discrimination during the pandemic

These innovations enabled us to share contextualized, granular data internally and to the public in an unprecedented way.

CCIS IN NUMBERS

- 26** reports released
- 1** Interactive dashboard
- 25** webinars recorded
- 5** live webinar events
- 12** PHC presentations
- 30+** data presentations (*to DPH staff, interagency workgroups, community groups, committees, advisory groups, national organizations, etc.*)
- 29K+** views on the CCIS website
- 25+** data to action discussions across DPH
- 7** DPH bureaus involved
- 60+** DPH staff volunteering to support
- 11** focus groups conducted
- 19** town/county level metrics released in tables
- 37** free text questions hand coded
- 39,206** free text responses hand coded
- 50+** CCIS Tweets from DPH Twitter
- 118** key findings released in our Q&A document

AVAILABLE CHAPTERS & RECORDED WEBINARS

Adult Survey

- General Methods/Descriptive Stats
- Personal Risk Mitigation
- Access to Testing
- Access to Healthcare
- Social Determinants of Health
- Vaccine Implications
- Mental Health
- Employment
- Substance Use
- Intimate Partner Violence
- Parents & Families

Youth Survey

- Part 1 - Mental health, changing responsibilities, and COVID experiences and perception
- Part 2 - Education and employment
- Part 3 - Youth safety and healthcare access

Population Spotlights

- Young parents
- Persons with disability
- Sexual orientation and gender identity (SOGI)
- Asian American and Pacific Islander (AAPI)
- Black
- American Indian and Alaskan Native (AI/AN)
- Hispanic/Latinx
- Discrimination/Framing Matters
- Housing
- Rural Communities
- Caregivers: adults with special needs and parents of children & youth with special healthcare needs
- Essential workers
- *Coming soon: Older Adults*

ACCESSING CCIS DATA / ANALYSIS

COVID-19 Community Impact Survey

The survey was done to better understand the many layered impacts of the pandemic. Please see below for a summary of key findings and other supporting materials.

TABLE OF CONTENTS

- Introduction
- Findings
- Background

HOW HAS COVID-19 IMPACTED YOU?



CCIS Data is posted on its own webpage:

- <https://www.mass.gov/covidsurvey>

On the website you will find:

1. Complete slide deck of all the CCIS data that has been released to date.
2. Recorded webinars for each of the chapters released to date
3. Data tables with CCIS data by MA county, municipality, and demographic groups.
4. Q&A document with more information about the survey and high-level talking points from each chapter and spotlight.
5. Interactive dashboard

These innovations also enabled us to
create critical change across the
Commonwealth



Population-focused

Prioritized inclusion of previously invisible populations

*48% of Parents of Youth
and Children with Special
Health needs reported
persistent poor mental
health (vs. 30% of other
parents)*

IMPACT

“We have been data poor, relying on limited data sources with small sample sizes. The kids and families we serve have great needs and have been historically unseen & unheard. The CCIS has provided a rich resource for us to make smart, strategic, evidence-based decisions that can make a difference in their lives.”

- CCIS Partner

Share needs of populations at state-level to inform policy making, in this case elevating need for respite care to be covered by Mass Health



Population-focused

Utilized data standards that are granular, inclusive, and reflective of populations' experiences

IMPACT

Mass Health is adopting the use of our rural definitions

BMC's Pediatric ED is utilizing CCIS SDOH questions

MAVEN now uses SOGI data standards

Codify equity in other data collection systems



**Prioritized community
engagement with
historically
marginalized
communities**

IMPACT

“Native Americans were once again visible in the data...The fact that CCIS connected the bureau with tribal members to pilot and then took their feedback and brought the data back was so important.”

- CCIS Partner

**Strengthened trust in DPH in communities
where there is a history of distrust**



Were nimble and shared breaking needs data for prioritization

CCIS illustrated the many unique barriers persons with disabilities face in accessing information and services related to COVID risk mitigation

IMPACT

“With CCIS data in mind, VEI prioritized improving vaccine access to people with disabilities. In the disability setting, people got vaccinated who wouldn’t have because of CCIS data.”

- CCIS Partner

Initiatives could quickly pivot to meet the needs of priority populations.



Utilized social justice framing when releasing results that drew linkages between inequities and systemic drivers

IMPACT

Health systems, municipalities and other entities conducting health needs assessments and improvement plans across the state, stated that the CCIS reports provided them with the evidence and framing needed to prioritize these systemic drivers in their health assessments and associated funding allocations.

Normalize the inclusion and naming of systemic drivers of inequities (structural racism, heterosexism, ableism) as health priorities

We didn't do everything right and learned
some lessons about where we can
improve

LESSONS LEARNED

Challenges

Time & human resource constraints

In roads with some priority populations

CBOs stretched thin

Distilling results into digestible formats

Being fully accountable to partners

Solutions

Building a team

Built relationships, will continue to do so

Provide financial support where possible

Building communications strategy

Consider full spectrum of work & engagement

We are now building on these lessons learned to create a sustainable data system that continues to engage communities, move our work upstream, and center health and racial equity.

VISION FOR CCIS 2.0



Replicate Successes and Incorporate
Lessons Learned from CCIS 1.0



Strengthen Relationships with Key
Partners Using Core Community
Engagement Principles



Build a sustainable data system
centered on racial and health equity
to identify and support policy and
practice ACTION



Reduce inequities in health
outcomes, including outcomes
experienced during the pandemic