

WELCOME!

While we are getting set up....

We would like to get to know our audience.



Please put your answer to the following questions in the chat.



Which organization are you affiliated with and what is your role?





The Consortium recognizes the uncertainty that ongoing changes at the federal level is causing for organizations working in public health and human service. The Consortium remains committed to its mission to reduce the human and economic burden of cancer in New York State (NYS).

It's important to remind ourselves that the Consortium's work is deeply rooted in science and data, and we are dedicated to implementing our mission and the New York State Comprehensive Cancer Control Plan with diligence and care.

The Consortium is also committed to its vision that people concerned about cancer will work collaboratively to implement the Cancer Plan while respecting and embracing the cultural, demographic, and geographic diversity within NYS. In fact, it's the collaborative and diverse nature of this voluntary organization that is our strength.

On behalf of the Consortium's Steering Committee, as New Yorkers let's continue to support and lean on each other.

Health Equity and Cancer: Navigating the Complexities of the Cancer Care Continuum

May 15, 2025

11 AM – 12 PM



NYSCC QUARTERLY MEETING SERIES



Housekeeping

Please mute your line.

If you have a question, please type it in the Chat Box.

Questions will be answered after the panel discussion.

This meeting is being recorded.

A link to the recording will be e-mailed to everyone who registered.

www.nyscancerconsortium.org





We are New Yorkers from all walks of life who work together to reduce the burden of cancer.

Member Area | New York State Cancer Consortium

NYSCC Quarterly Meeting Series: Health Equity and Cancer

Upcoming Meetings

- Survivorship (Part 1)
 Tuesday, September 16th, 11:00 AM to 12:00 PM
- Survivorship (Part 2)
 TBD

Events | New York State Cancer Consortium



Pre-Meeting Poll



Health Equity and Cancer:

Navigating the Complexities of
the Cancer Care Continuum



Session Objective:

After participating in this session, participants will understand successful strategies that influence the development of inclusive solutions in cancer care, the crucial role of the Consortium in fostering collaboration among stakeholders and supporting research that emphasizes diversity. Our ultimate goal is to ensure equitable access to cancer resources for all.

Meet Our Speakers



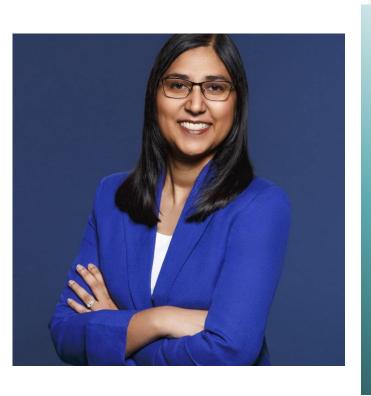


Alyson Moadel-Robblee, PhD

Health Psychologist, Professor of Clinical Epidemiology, and Founding Director of the BOLD Cancer Wellness Program

Montefiore Einstein Comprehensive Cancer Center (MECCC)

Dr. Moadel-Robblee is also Deputy Director of Community Outreach & Engagement at MECCC, where her charge is to ensure our cancer center's clinical and scientific efforts to promote equitable and optimal cancer care from screening and clinical trials to survivorship are informed by the patient and residential community it serves. A major focus of her programmatic and scientific work is on the use of "extender" patient navigators, including peer and virtual, in-patient engagement efforts.



Dr. Pia Banerjee

Director, Cancer Innovation & Transformation

American Cancer Society (ACS)

As a globally recognized researcher, clinical neuropsychologist, and health technology executive, Dr. Banerjee has built and scaled digital health tools that impact tens of millions of individuals annually.

Before joining ACS, she served as Senior Vice President at Neuroglee Therapeutics, where she led worldwide teams in developing Al-enabled health solutions for patients, caregivers, and providers.

Dr. Banerjee's diverse career across industry, academia, and clinical care has directly shaped global clinical standards, with her research at St. Jude Children's Research Hospital recognized as one of the most impactful scientific achievements of the year.

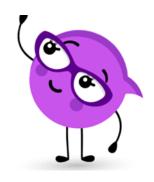
Dr. Banerjee completed her undergraduate degree at MIT, her Master's degree and PhD in Clinical Psychology at Washington University in St. Louis, and her postdoctoral fellowship at UCLA.

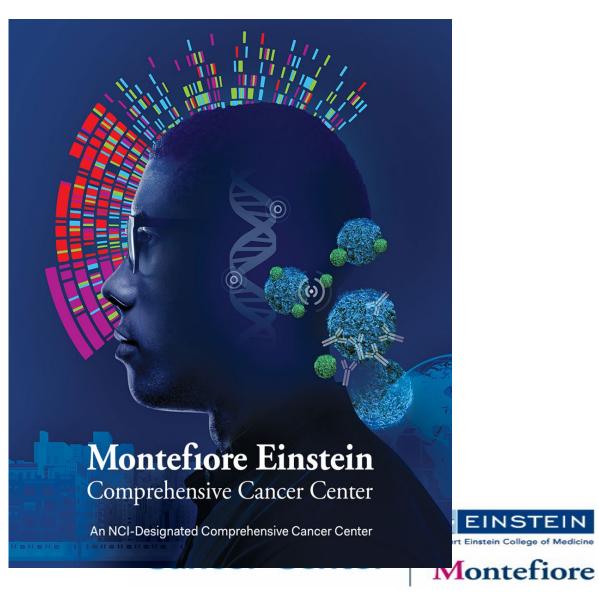


Widening the Lens on Patient Navigation: Peer and Virtual Models In Action

Alyson Moadel-Robblee, Ph.D.
Founding Director,
BOLD Cancer Wellness Program
Deputy Director,
Community Outreach & Engagement
NYSCC, Steering Committee Member







Overview

- Describe a Peer Navigation Model that developed out of a cancer patient needs assessment
 - Need, Impact, and Sustainability
- Describe an Al-driven Patient Navigation initiative
 - Conception, Impact, Future Directions

Setting: Bronx, NY

Population: 1.4 million

Poverty: 31%

Race/Ethnicity:

• Hispanic 57%

• Black 44%

Foreign born: 34%

Common Ancestries:

- Dominican (22%)
- Puerto Rican (20%)
- West Indian (8%)
- Mexican (6%)
- Jamaica (5%)
- Sub-Saharan African (5%)

Non-English as primary lang: 56% Single-person households: 32%



U.S. Census Bureau Quickfacts: Bronx County, New York, www.census.gov/quickfacts/fact/table/bronxcountynewyork/PST045223. Last updated in 2023 Accessed 23 May 2024.

Statistical Atlas, https://statisticalatlas.com/county/New-York/Bronx-County/Ancestry, Last updated in Sep 4, 2018. Accessed 23 May 2024.

Cancer Burden & Disparities

High Incidence, late stage diagnosis, and/or mortality:

- > Prostate
- Colorectal

> Breast

Cervical

➤ Lung

> Multiple Myeloma

> Liver

➤ AML/MDS (blood cancers)



Risk Factors

- Physical
 - Pain/fatigue
 - Disfigurement/Disability
- Emotional
 - Depression/Anxiety
 - Re-traumatization
 - Body image distress

- Social
 - Loss of roles/agency
 - Loneliness
 - Stigma
 - Medical mistrust
 - Financial toxicity



Risk Factors for Increased Cancer Burden

Bronx is # 62 of 62 counties in health factors including:

- Obesity
- Asthma
- Food insecurity/houselessness
- Population density
- Air pollution
- Crime
- Limited green space
- Lower physical activity





31% living at poverty line



"There is no power for change greater than a community discovering what it cares about"

Margaret J. Wheatley

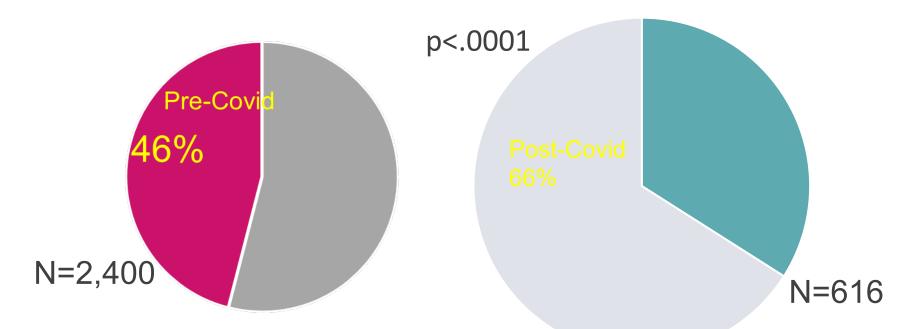






High levels of Distress reported by Bronx Cancer Patients (Pre/Post-COVID)

Clinical Distress > 4



*Previous studies have found an average prevalence of 39% of clinical distress based on the DT

Mitchell AJ, JCO, 2007

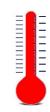


Contributors to Distress (P<.001)

	High Distress	Low Distress
Housing/Insurance/	24%-36%	11%-15%
Transportation		
Partner/Children	18%	7 %
Worry/Sadness/Depression/Nervous	67%-83%	16%-32%
Spiritual Concerns	13%	6%
Pain/Fatigue	65%-70%	32%-34%
Sleep problems	67%	31%
Tingling in hands/feet	50%	33%



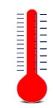
Medical Mistrust



- People of my ethnic group receive the same medical care from doctors and HCWs as people from other groups
 - > 20% Disagree
 - > 22% Neutral

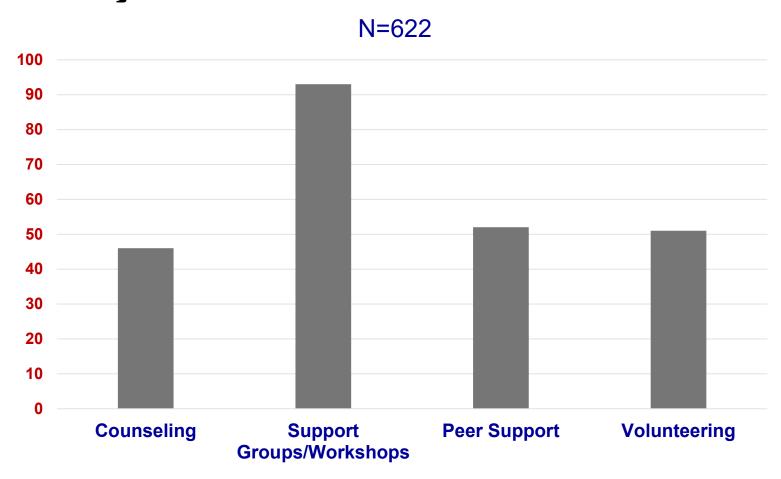
- I have personally been treated poorly or unfairly by doctors or HCWs because of my ethnicity
 - > 12% Agree
 - > 9% Neutral

Isolation/Loneliness



- live alone
 - > 22%
- "I miss having people around"
 - > 42%
- "There are many people I can rely on when I have problems
 - > 14% No

Psychosocial Needs Endorsed



BOLD WELLNESS WORKSHOPS

These workshops aim to foster a sense of community and connection. Held at the newly opened BOLD Cancer Wellness Center in the Bronx, we offer workshops on a weekly or monthly basis. Some popular workshops include:

CREATIVE ARTS

- Paint Club
- Conversational Spanish
- Telling Your Life Story (Writing workshop)

Services may

be virtual.

in-person or

hybrid!

- Crochet
- Book Club

MIND/BODY WORKSHOPS

- Yoga
- Meditation
- Reiki
- Stress Management
- Spiritual Infusion Sessions

BOLD WELLNESS EVENTS

Throughout the year, we host beauty days, games, giveaways, raffles, food, and fun to celebrate life, love, holidays, and self-care.

NUTRITION/PHYSICAL ACTIVITY

- Diabetes Prevention Education
- Dance/Fitness
- Nutrition/Cooking Class

EDUCATIONAL PROGRAMS

 Our educational programs cover a wide range of topics, including Body Image, Relationship/Intimacy, Returning to Work, Life After Cancer, Spirituality, Managing Treatment Side Effects, Communicating Your Needs, and more.

What is BOLD?

BOLD Living is an integral part of Montefiore Einstein Comprehensive Cancer Center's dynamic Community Outreach and Engagement program. This psychosocial and Integrative Oncology initiative arose from patient and community input, highlighting the demand for an outpatient wellness and support program accessible to all those affected by cancer in the Bronx. This encompasses Montefiore patients, family members, caregivers, and individuals under care elsewhere. Everyone is invited, and all services are offered free of charge.

Our dedicated psychosocial team collaborates closely with the primary oncology team and other sub-specialties as necessary to design services and connections that address the emotional, physical, and spiritual needs of our patients and the broader cancer community.

Through a multidisciplinary approach, mindbody practitioners, mental health specialists, trained interns, and cancer survivor volunteers unite to offer a comprehensive and inclusive array of services in both English and Spanish.





Ponce Bank

Center of Excellence

BRONX
ONCOLOGY
LIVING
DAILY

A CANCER WELLNESS
INITIATIVE FROM THE
PSYCHOSOCIAL AND
INTEGRATIVE ONCOLOGY
PROGRAM

WHERE MIND, BODY, AND SPIRIT ARE NURTURED.

Montefiore Einstein Comprehensive Cancer Center

BOLD Buddies – Peer Navigators

 The B.O.L.D. Buddies provide peer support, treatment companionship, cancer experience navigation, shared understanding, inpatient bedside visits, and community outreach support





Impact of BOLD engagement on Loneliness in 67 breast cancer patients p=0.017

Did the patient engage in services?	N	Mean Loneliness change score over 3 mos	Standard Deviation	Standard Error Mean
No	34 (~50%)	0.5294	2.64272	0.45322
Yes	33 (~50%)	9697	2.35166	0.40937

 Those who did not engage in BOLD demonstrated a 9% increase in loneliness, those who did engage saw an average 16% decrease in loneliness scores.

Impact of Peer Navigation

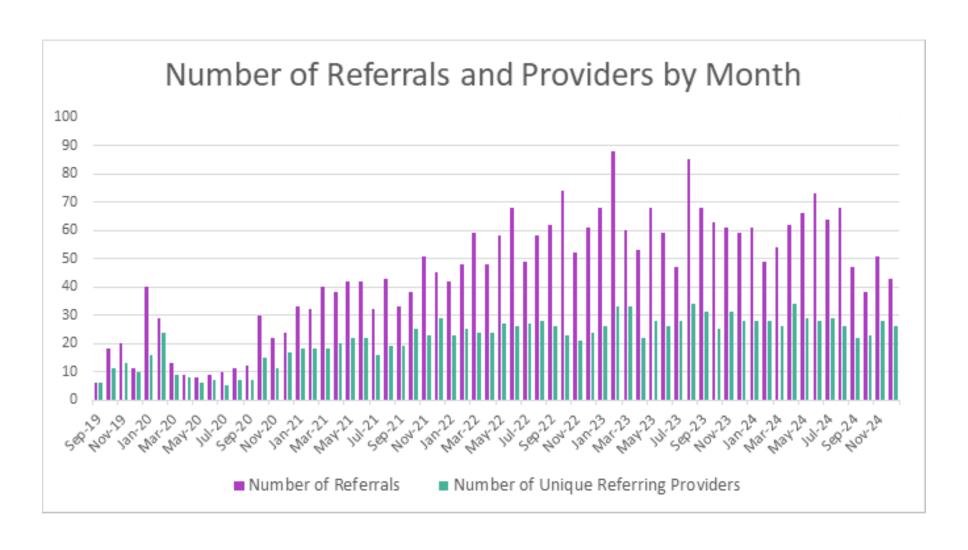
- NEED: 47% of patients express interest in a BOLD Buddy
- REACH: 301 cancer patients over pandemic
- <u>SATISFACTION</u>: 89% of patients rate Buddy support as very much to extremely beneficial to their care

Key areas of benefit (endorsed by patient):

- 81% Buddy helps in <u>reducing distress and isolation</u>
- 63% help in communicating with health care team about treatment side effects and medical issues
- 57% helps in trusting medical care & recommended treatments
- 46% helps in adhering to appointments & treatment

Peer navigation in EPIC workflow

(Sept 2019 - Dec 2024)



Overview

- Describe an Al-driven Patient Navigation initiative
 - Conception, Impact, Future Directions







•<u>Alyson B. Moadel et al.</u> Al virtual patient navigation to promote re-engagement of U.S. inner city patients nonadherent with colonoscopy appointments: A quality improvement initiative. Presented at ASCO, 2024.

Challenge:

From 2022-2023, 20% of Bronx patients did not complete scheduled colonoscopy (2,400 vs 8,783)



QI Project: Colonoscopy Re-engagement Initiative

- Outreach is typically one-way communication
 - Directive, Prescriptive, Navigational
 - Limitations to outreach: reduced buy-in => non-adherence
- Engagement is bi-directional communication
 - Educating, Motivating, Discussing, Addressing barriers to care
 - Limitations to Engagement: human resources
- Opportunity: layered/paired navigation
 - e.g., MyEleanor





 Employed MyEleanor between Apr-Dec 2023 to target reengagement of 2,400 of 11,183 English- and Spanish-speaking patients nonadherent with colonoscopy appointments in 2022-2023.

MyEleanor, an English/Spanish speaking Al virtual navigator:

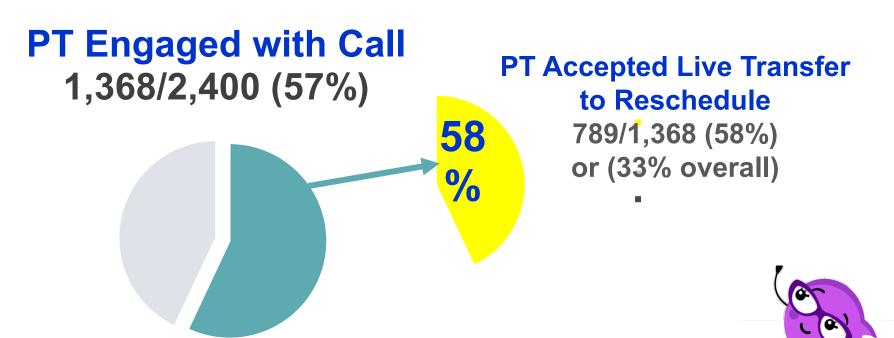
 (a) called patients to invite rescheduling, (b) assessed barriers to uptake, c) offered live transfers to clinical staff to reschedule, and d) provided procedure prep reminder calls.

Evaluable outcomes:

- (a) engagement with MyEleanor, (b) live transfers accepted,
 - (c) colonoscopy completion rate, and (d) patient volume, with
- (e) barriers to care, and (f) predictors of actionable engagement examined.



Results: Engagement with MyEleanor



Results: Engagement with MyEleanor

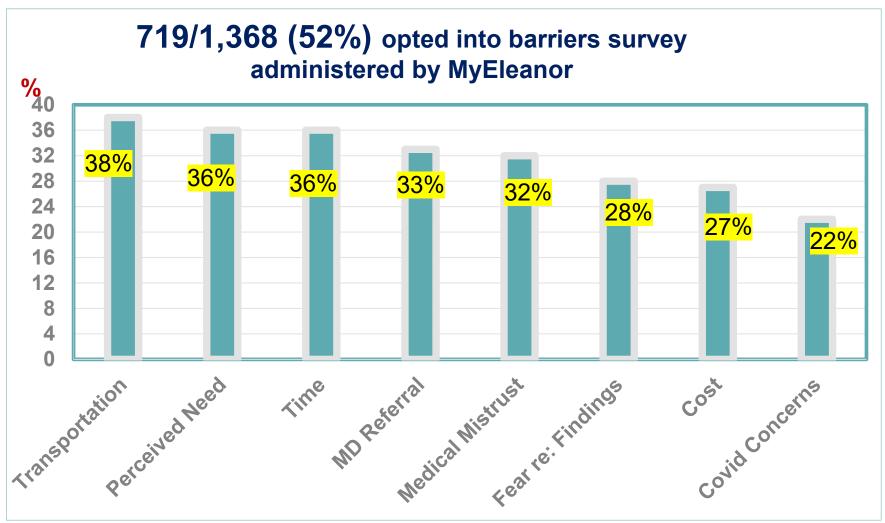


- Patients who accepted transfer were 25% more likely to complete colonoscopy
- No-show completion rate nearly doubled from 10% to 19%
- Sociodemographic predictors of engagement:
 - Patients who engaged were a Mean age of 56.66 (41-79 yrs), female (66%), Hispanic (41%), Black (33%), English (75%) or Spanish (25%) speaking, and single/D/W (63%), and Unemployed/Disabled (49%)

Those unemployed/disabled accepted transfer more often (49%) than those employed/retired (41%), chisquare=6.70(4),p<0.02.

Barriers to Screening





Barriers: Predictors of Engagement/Action

Accepting transfer:

More likely:

Greater # of barriers

Less likely:

- Cost
- Medical Mistrust
- Cultural concerns

Completing procedure:

Less likely:

- Cost
- Fear (procedure/findings)

Spanish-dominant patients and those **declining to identify their race** reported nearly twice the number of barriers, F(599)=47.48 and F(571)=56.66, p<0.001, respectively.

Volume Increase



- Patient volume increased by 36% (1,363 patients)
- Completed colonoscopies:

2022: 3,8982023: 5,261

- 52 hours month freed up for 7 Patient Navigators to:
 - Schedule new patients
 - Field MyEleanor live transfer calls

OUTREACH AND ENGAGEMENT in action

Take Home Points - Peer Navigation

Benefits and Outcomes

- Trusted guide to the health care system
- Role modeling of health activation
- Shared lived experience & Advocacy
- High patient interest for PN (50%)
- Enhances trust in medical system/providers
- Enhances self-reported medical adherence
- Reduces loneliness & distress

Sustainability:

- Hospital Volunteer Dept offer infrastructure
- 1:1 ratio of PN need and volunteer interest
- Integration into EHR referral = "standard care"
- Ongoing marketing and research for visibility, growth, and impact



Take Home Points – Virtual Navigation

Al-driven virtual navigation demonstrated:

- feasibility through high patient acceptance and engagemer
- clinically significant impact on patient re-engagement with
- Increased patient volume through increasing capacity of human patient navigators
- Ability to identify barriers to care among a substantial subgroup of patients

Next Steps:

 Measuring MyEleanor's impact on PN burden/burnout, other screening programs (breast/lung), distress/SDOH screening, cost savings, stage shifting

Challenges:

 Integration of an outside technology application with hospital-based EMR system/tracking system – extensive data mining and synthesis efforts required! It takes a Village





ACS ACTSTM

Access to Clinical Trials & Support





The Problem: Barriers to Clinical Trials

- Only 7% of cancer patients participate in clinical trials, with participation from historically underrepresented communities at just 4%
- 20% of cancer trials fail due to insufficient enrollment

Administrative Burden

Medical Records

Trial Match Challenges

- Lack of onsite trials
- Complex eligibility criteria

Limited Awareness

- Provider lacks time to discuss/research trials
- Unconscious biases



Financial Burden

- Transportation & lodging
- Lost wages

Health-Related Social Needs

- Food insecurity
- Affordable housing & transportation



Our Solution





ACS ACTS™: Access to Clinical Trials & Support

- The American Cancer Society brings you ACS ACTS to connect you with personalized clinical trial options and the right resources by your side.
- In collaboration with Massive Bio, ACS reduces the common barriers to clinical trial enrollment and participation.
- Initial launch date: February 24, 2025



ACS ACTS

Strategic Goal:

Improve equitable access to cancer clinical trials by reducing the barriers to enrollment and participation

Key Constituents:

Patients, healthcare providers, and caregivers

Program Offerings:





Program Eligibility

Age

All ages

Cancer Types

All cancer types

Location

- Feb 2025: Regional launch¹ for patients living in or willing to travel to Northeast US²
- Fall 2025: National launch expected; no location restriction

Interest

Interested in exploring eligible clinical trials:

- All interventional cancer clinical trials listed on clinicaltrials.gov
- Agnostic to sponsor

¹The Northeast region contains the largest concentration of clinical trial sites in the US and includes regions where ACS can more easily provide transportation and lodging for clinical trial participation.

² Connecticut, Delaware, Maine, Maryland, Massachusetts, New Hampshire, New Jersey, New York, Ohio, Pennsylvania, Rhode Island, Virginia, Vermont, West Virginia, or Washington, D.C.



Program Costs

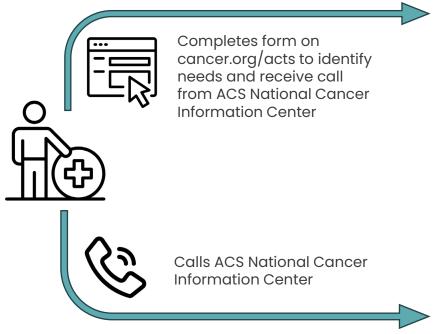
ACS ACTS is **FREE!**

ACS is a non-profit, with programming supported by donations and sponsorships

Some ACS programs for health-related social needs involve a small cost, which the patient is informed of (e.g., prosthetics)

Workflow Part 1: Intake to Trial Matching







- Addresses health-related social needs
- Provides clinical trials education
- Assesses eligibility for ACTS
- Makes warm handoff on 3-way call via vanity line to Massive Bio if constituent is interested



- Collects consents
- · Obtains medical records
- · Performs Al-based clinical trials matching
- Emails matches with phone follow-up, and SMS or email follow-up as backup
- Notifies treating provider of match results if patient consents
- Encourages patient to discuss with treating provider and can arrange 3-way call; or patient can talk to Massive Bio physician

Patient, Caregiver, or Healthcare Provider

ACS

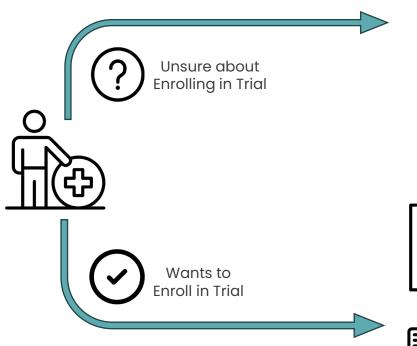
National Cancer Information Center Team

MassiveBio

Oncology Nurses & Patient Coordinators

Workflow Part 2: Matching to Needs Met







Follows up with patient at patient's desired intervals

If unable to make contact, receives automated follow-up questionnaire via SMS/email to track interest



Coordinates with trial site; tracks referral and enrollment via SMS/email questionnaire



Conducts needs assessment by phone:

- 1) Before trial enrollment
- 2) 1 Month post-enrollment
- 3) Monthly if requested

If needs identified, calls NCIC with patient via vanity line

Patient

MassiveBio

Oncology Nurses & Patient Coordinators



Addresses identified needs by coordinating ACS services:

- Lodging
 - Extended Stay America
 - Hope Lodge (via HCP)
- Financial
 - Patient Advocate Foundation
- Transportation
 - Road to Recovery
- Prosthetics & Survivor Care Supplies
 - Ever You
- Social Support
 - ACS CARES app
 - · Cancer Survivors Network
- Local & National Resources
 - FindHelp.org



CoC Accreditation: Standard 9.1



CoC Standard 9.1 requires accredited cancer programs to enroll a specified percentage of eligible patients into cancer-related clinical research studies.

ACS ACTS can help fulfill compliance with CoC Standard 9.1



To count an ACS ACTS case toward CoC Standard 9.1, providers should follow CoC requirements and their internal procedures, which may include:

- Initiate and document referral to a qualifying trial prior to enrollment
- Maintain written verification of enrollment

If provider engages with ACS ACTS on behalf of patient:

Considered a provider referral; counts if CoC requirements are followed

If patient or caregiver initiates ACS ACTS participation:

- MassiveBio will notify provider of trial matches prior to enrollment if patient consents, and encourage patient to discuss trial match results with provider
- Self-referral becomes a provider referral; counts if CoC requirements are followed

Note: This is a general informational guide. Follow your institution's internal procedures and the Commission on Cancer Standards.



Metrics: 2/24/25 - 4/29/25

Program Participants

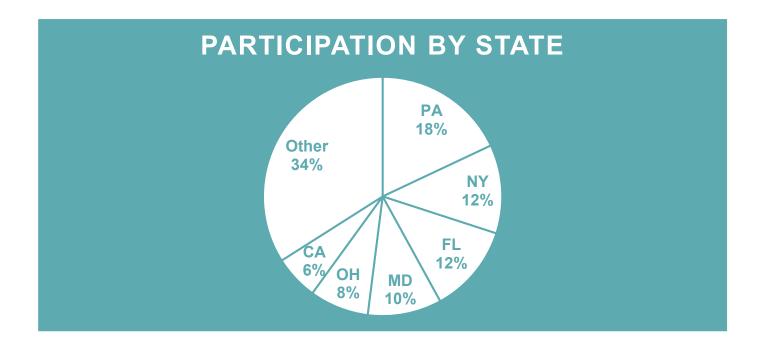
314

Percentage with Health-Related Social Needs

99%

Number of Health-Related Social Needs Identified

831



ACS ACTS cancer.org/acts

American Cancer Society ACTS

Sign up for ACS ACTS today! Scan the QR code to fill out the form or call our 24/7 National Cancer Information Center line at 1-800-227-2345.





For more information on ACS ACTS, go to cancer.org/acts or reach out to Dr. Pia Banerjee: pia.banerjee@cancer.org

Post-Meeting Poll



Question & Answer



Join the Consortium! and look out for upcoming meetings



Events | New York State Cancer Consortium

Thank you for Attending



cancerconsortium@health.ny.gov



