

## OBJECTIVES

### PURPOSE

- These clinical trials aim to answer critical questions for cancer treatment in clinical practice and promote inclusivity in cancer research.
- The selected trials span liquid and solid cancer, including treatment, radiation, and imaging theranostics.
- These trials are typically not sponsored by industry but answer important practice questions on how to treat patients to improve patient outcomes.
- These practice changing trials will inform survival, how to improve toxicity, identify best current treatment for disease, and the best way to deliver therapy.

## METHODS

### INNOVATION

In 2023, four investigator-initiated trials written with focus on community engagement and inclusion.

- Clinical trials are written by the physician investigators
- Various decentralized tools including:
  - Real world data from the EHR to correlate treatments
  - Virtual consent using digital signature capture
  - Virtual Data capture
  - Using imaging to deescalate cancer treatment to minimize patient burden.

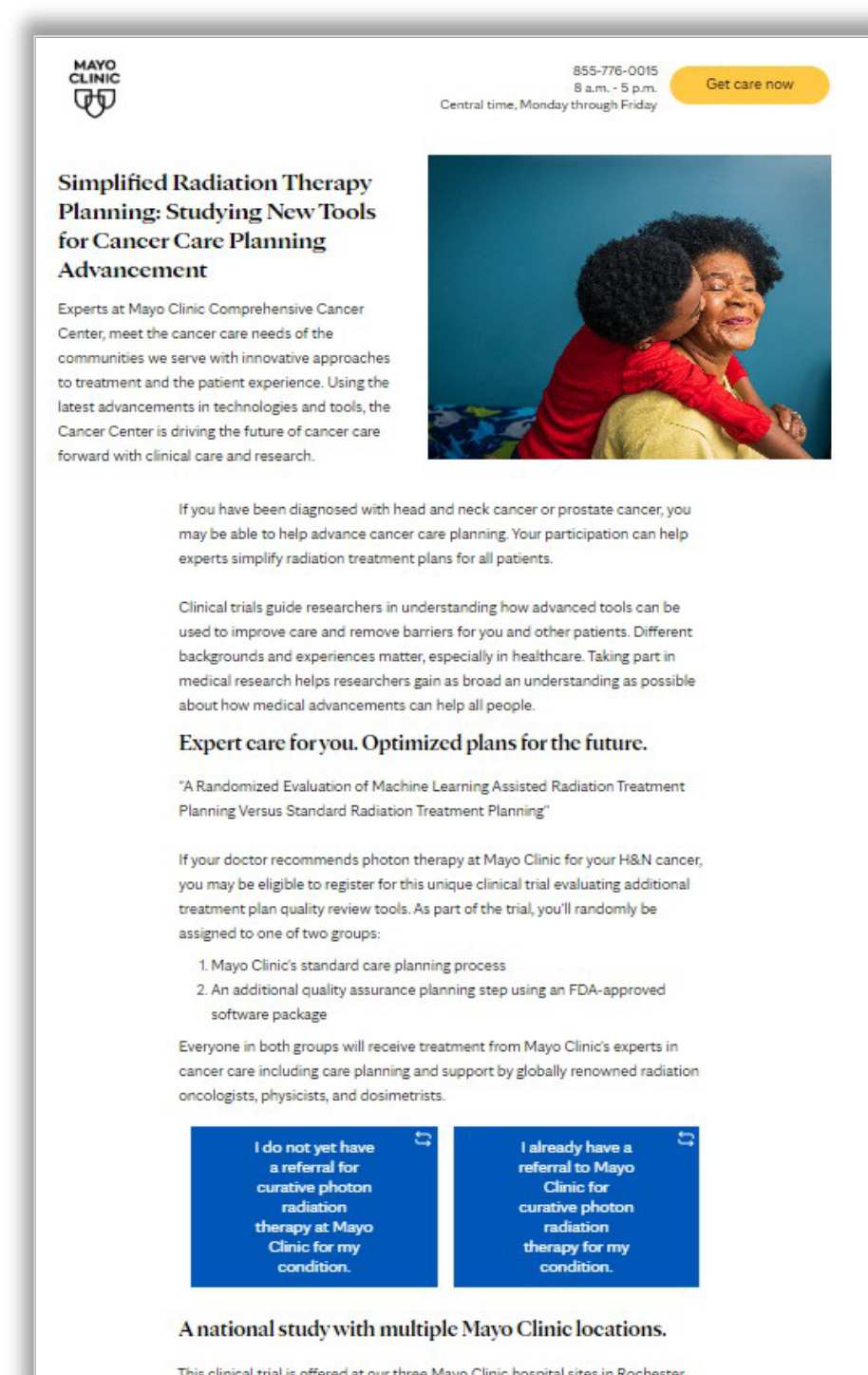
### RECRUITMENT PLAN

Deployed recruitment techniques include:

- Marketing in print and social media
- Development of study specific websites
- QR codes to link to the study site and consent information
- Partnership with national organizations
- Consultation with community partners for further consideration to meet inclusive needs

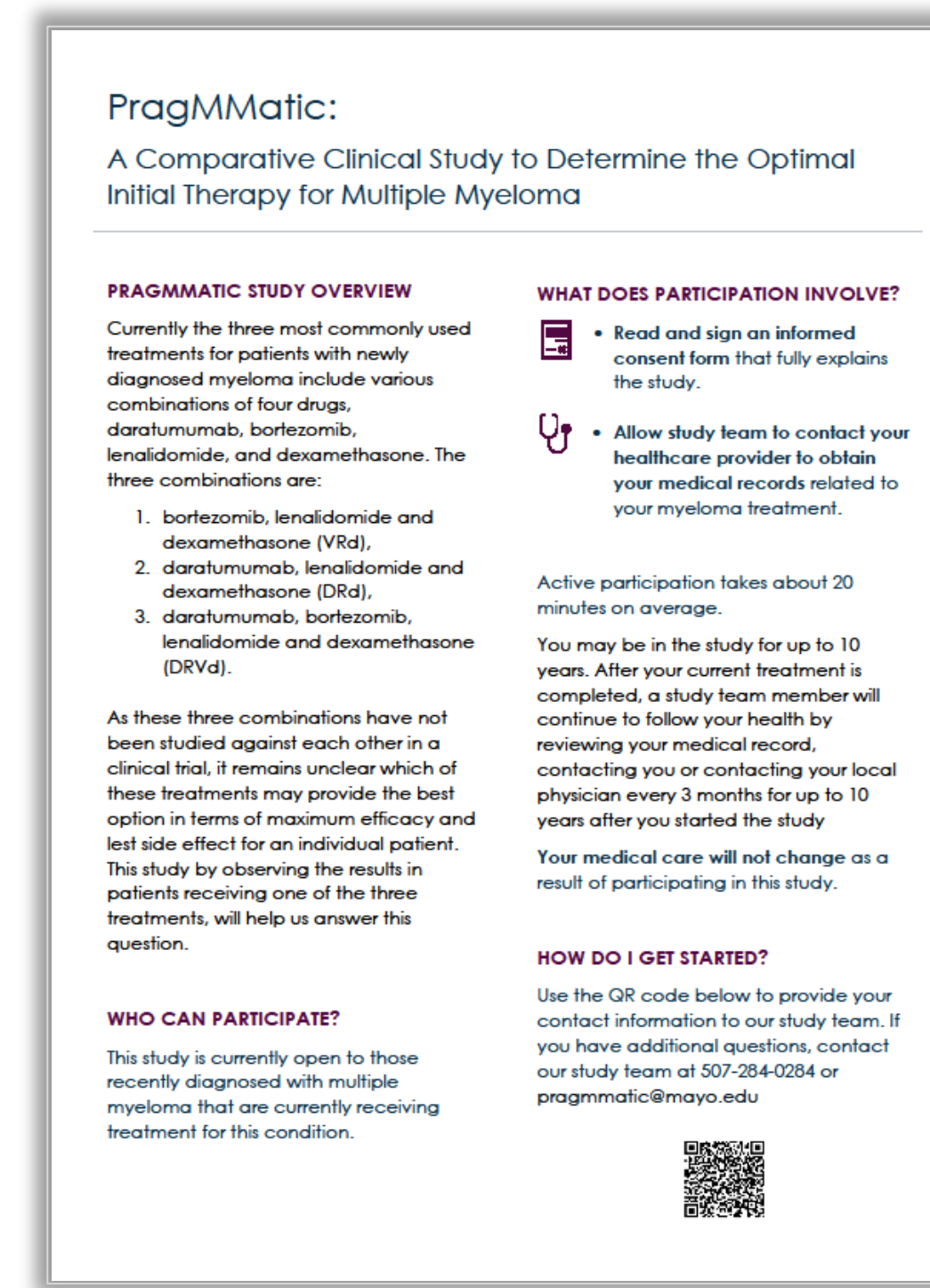
## RESULTS

FIGURE 1.



Website example

FIGURE 2.



Trial Flyer Example

TABLE 1: Investigator-Initiated Trials

Trial	Purpose
Trial 1: NCT60151717	What is the BEST initial therapy for multiple myeloma
Trial 2: NCT05979883	Determine if machine learning assisted radiation treatment planning improves treatment versus standard radiation treatment planning
Trial 3: NCT06037863	Is there an impact on radiation treatment when a bladder is full versus empty.
Trial 4: NCT06200103	Can post therapy imaging be used to personalize and optimize the use of radionuclide therapy for prostate cancer.

Four Investigator-Initiated Trials with focus on community engagement and inclusion

## DISCUSSION AND CONCLUSIONS

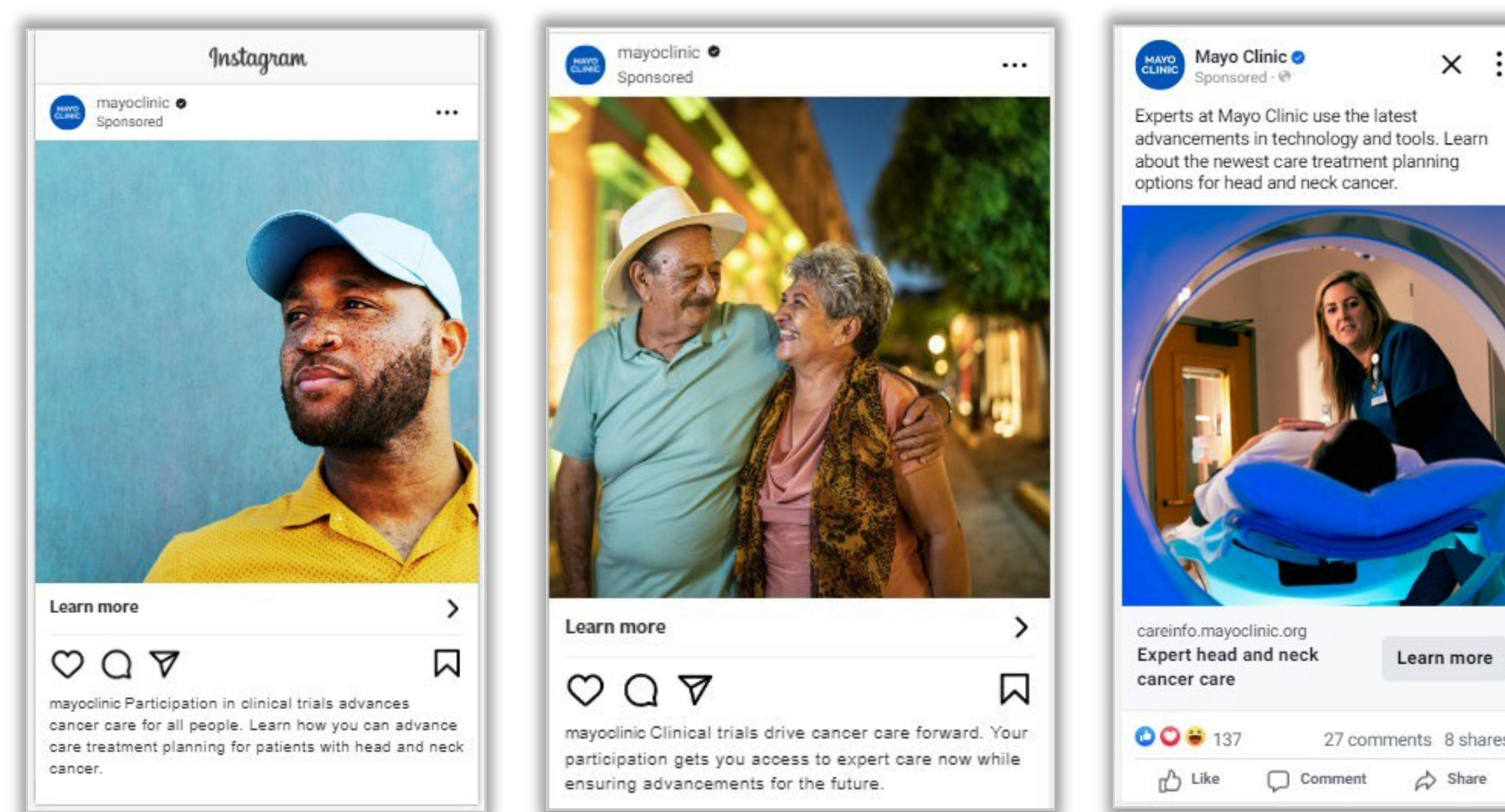
### TIMELINES

- Trials were written, activated, and opened to accrual in 3-5 months.
- Accrual is currently open throughout the 4 states that the multi-site center spans.
- National accrual is open for NCT60151717.
- These inclusive designs and decentralized tools permitted:
  - Greater participation of underrepresented patient populations.
  - Developed processes and technology to support clinical trials.

### WHY DO THESE TRIALS MAKE A DIFFERENCE?

- Each trial seeks to answer questions that will directly influence cancer care.
- The inclusive design of each protocol seeks to assure broad representation that will allow better understanding of how treatments affect individuals.
- Through inclusive designs and broad eligibility criteria, the trials are expected to accrue faster and inform standard of care.

FIGURES 3-5.



Figures 3-5 are sample Clinical Trial ads.