

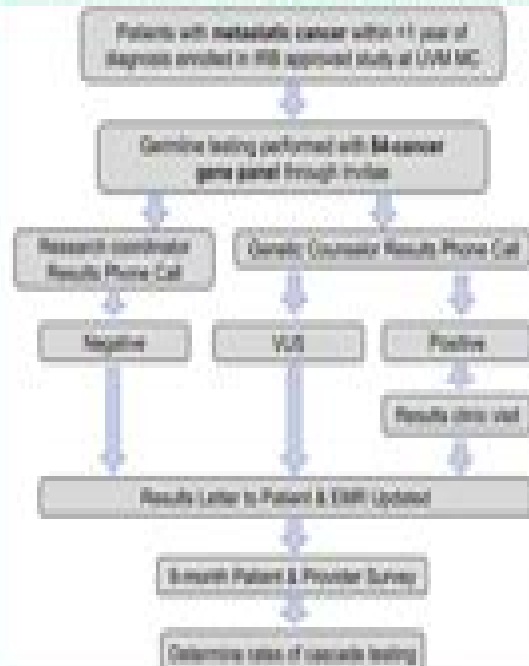
ELIMINATING BARRIERS TO UPTAKE OF GERMLINE GENETIC TESTING FOR PATIENTS WITH ADVANCED CANCER: THE GTAC STUDY

Kara E. Landry, MD, Stephen J. Foley¹, Heather M. Feldman, PhD, Sarah A. Novak, PhD, Hannah L. Taylor², Marc R. John³,
Wendy C. McKinnon, MD, Laura K. Finkel, MD, Sarah M. Nelson, MD, Edward B. Espino, MD, PhD, Marc A. Greenblatt, MD, Marc E. Reed, MD⁴
¹Division of Oncology and Hematology, University of Vermont Medical Center, Burlington, VT; ²Center for Health Systems Research and Analysis, University of Vermont, Burlington, VT
³Department of Pathology & Laboratory Medicine, UVM Medical Center, Burlington, VT; ⁴Center for Cancer Control, University of Vermont, VT

Introduction

- Identification of pathogenic germline variants (PGVs) is important for patients with advanced cancer: guiding treatment decisions, cancer screening, and testing of family members.
- Despite this, barriers to testing still exist.
- Streamlined processes for genetic testing are needed.
- We assessed the feasibility of a novel genetic counseling/testing approach:
 - Genetic testing with written education was performed.
 - Genetic counseling *only* for those testing positive.

Methods



Results

Figure 1. Enrollment

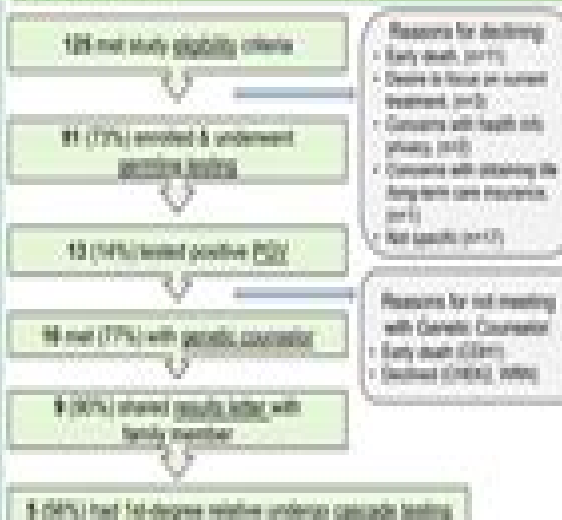


Table 1. Participant Characteristics

Total Participants, n	81
Age (years), mean ± SD	66 ± 12.8
Sex	42 (51.7%)
Caucasian, non-Hispanic	64 (79.0%)
Rural residence	43 (53.1%)
Cancer types, n (%)	
Colorectal	27 (33.3%)
Lung	27 (33.3%)
Prostate	19 (23.4%)
Rectal	4 (4.9%)
Breast	3 (3.7%)
Melanoma	3 (3.7%)
Brain	3 (3.7%)
Sarcoma	3 (3.7%)
Other	7 (8.6%)

Table 2. PGVs identified

Total, n (%)	13 (16.0%)
BRCA1	3 (38.5%)
BRCA2	3 (38.5%)
CHD8	2 (25.6%)
MTOR	1 (12.8%)
MLH1	1 (12.8%)
MSH2	1 (12.8%)
CDKN2A	1 (12.8%)
APC	1 (12.8%)

*Genotype tested for BRCA1 and BRCA2 only. Germline testing performed through Invitae.

Table 3. Satisfaction Survey Responses

Patient Survey Responses, n=74 (91.2%)	Strongly Agree	Agree
Satisfaction with testing process		
I am pleased I had the genetic test	94.6%	
I preferred having testing incorporated to existing visit	94.6%	
Effectiveness of patient education with testing results		
I understood the implications of results for me	96.1%	
I felt confident in my ability to explain my test results	92.0%	
I felt confident in my ability to explain what my test results mean to me	96.1%	
I felt confident in my ability to explain what my test results mean for others	92.0%	
Medical Oncologist Survey Responses, n=18		
Satisfaction with testing process	100%	
Genetic testing approach was easier than the previous process	83.3%	
Genetic testing results were obtained in broader fashion	83.3%	
Preferred testing first and counseling only patients with positive results	100%	
Patient impact on clinical practice		
Test results helped make treatment decisions	100%	
Increased willingness to order genetic testing for patients	88.9%	
Felt comfortable recommending testing initiation of patients with positive results	100%	
Continued interest in testing process		
As an option for patients with metastatic cancer	88.9%	
Expanding offer to all patients with cancer	83.3%	

Conclusions

- This paradigm shifting process of "testing first and counseling after" is feasible and acceptable for cancer patients.
- Patients and providers report high satisfaction with this process that removes known barriers to testing, such as extra clinic visits.
- Increased rates of cascade testing were clinically meaningful compared to current published rates.
- Further use of this process will require delivering education to patients prior to testing and ensuring that providers are comfortable ordering testing.

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