Biomarker testing patterns among patients newly diagnosed with prostate cancer in a community oncology network

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Key Takeaway

Biomarker testing patterns and time-to-treatment initiation varied widely among patients newly diagnosed with prostate cancer in the community setting, where most patients receive their cancer care

Conclusions

The study findings provide insights into the frequency of biomarker testing among patients newly diagnosed with prostate cancer receiving care in a large community setting



Despite guideline recommendations for biomarker testing, only a small percentage of patients overall received it, with slightly higher rates among those diagnosed with stage IV cancer



To realize the potential benefits of targeted therapies, there is a need to improve biomarker testing rates to better determine which patients are eligible for targeted treatment



The findings suggest that opportunities exist to increase biomarker testing education for healthcare providers in the community setting and to design tailored interventions to increase testing among patients with prostate cancer

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Link: https://www.congresshub.com/Oncology/GU2025/GeneralProstateCancer/Mahmud

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Background

ickground			
Prostate cancer guidelines recommend that clinicians order biomarker testing for patients with metastatic prostate cancer ¹ Tumor biomarker testing is performed to identify potential actionable mutations that can inform appropriate treatment strategies as well as determine a patient's eligibility for clinical trial participation The study aimed to evaluate biomarker testing patterns among US patients newly diagnosed with prostate cancer who received			ent strategies •
are in the community setting	parterns among oo pa	licites newly diagnosed with prostate cancer w	•
esults			
The cohort consisted of 18,706 patients aged ≥	18 years who were new	ly diagnosed with prostate cancer	
/lean age at diagnosis was 71.0 years (standard Black or African American, and 0.8% were Asiar		1% of patients were non-Hispanic White, 12.6%	% were Hispanic, 7.1% were
he primary insurance coverage for the majorit	y of patients was comn	nercial (59.8%), followed by Medicare/Medicai	d (39.8%)
ewer patients were diagnosed in early (I and II)) than in late (III and IV)	stages of the disease, 32.4% and 38.3%, respe	ectively
TABLE 1: Patient demographics and clinical			
Characteristic	N=18,706	Characteristic	N=18,706
Mean age at diagnosis, years (SD)	71.0 (9.1)	ECOG PS score, n (%)	
Race/ethnicity, n (%)	2262 (126)		10,611 (56.7) 2124 (16.7)
Hispanic ^a Non-Hispanic Asian	2362 (12.6) 152 (0.8)	2–5	3124 (16.7) 543 (2.9)
Non-Hispanic Asian Non-Hispanic Black or African American	132 (0.8)	Z=0 Missing	4428 (23.7)
Non-Hispanic Other ^b	4794 (25.6)	Karnofsky Performance Scale score, n (
Non-Hispanic White	8670 (46.4)	90–100	10,992 (58.8)
Unknown/declined to inform	1397 (7.5)	70-80	2842 (15.2)
Provider location ^c , n (%)		0-60	444 (2.4)
Rural	92 (0.5)	Missing	4427 (23.7)
Urban	18,614 (99.5)	Histopathology, n (%)	
Primary insurance coverage, n (%)		Adenocarcinoma	2489 (13.3)
Commercial	11,185 (59.8)	Small cell carcinoma	29 (0.2)
Medicare/Medicaid	7445 (39.8)	Squamous cell carcinoma	2 (0.01)
Unidentified or self-pay	76 (0.4)	Missing	16,186 (86.5)
Disease stage at diagnosis, n (%)		Comorbidities ^d , n (%)	
	1464 (7.8)	Other cancer	2568 (13.7)
	4594 (24.6)	Hypertension	2332 (12.5)
	2656 (14.2)	Nutrition deficiency	1454 (7.8)
IV	4510 (24.1)	Hyperlipidemia	1269 (6.8)
Missing	5482 (29.3)	Diabetes	938 (5.0)

ludes Hispanic or Latino American Indian/Alaska Native. Hispanic Asian. Hispanic Black or African American, Hispanic Native Hawaiian/Other Pacific Islander, Hispanic other, and Hispanic White t excludes "unknown/declined to inform").

spanic multi-race, and Non-Hispanic Native Hawaiian/Other Pacific Islander les Non-Hispanic American Indian/Alaska Native, Non-His

ovider location for two respondents was unknown. ^dComorbidities occurring in \geq 5% of patients.

ECOG PS, Eastern Cooperative Oncology Group performance status; SD, standard deviation.

Reference **1.** NCCN Clinical Practice Guidelines in Oncology. Prostate cancer. Version 4.2024; May 17, 2024.

Methods

- etrospective database analysis of patients receiving care in a large oncology practice in the West South Central region of the United States was conducted
- roximately 1.7 million unique patients with various cancer types
- e analysis included patients newly diagnosed with prostate cancer between January 1, 2018, and June 30, 2022 ng the study period
 - Biomarker testing was performed for 2548 (13.6%) of patients with prostate cancer. Among all patients diagnosed with stage IV prostate cancer (n=4510), 1044 (23.2%) received biomarker testing
 - The mean (SD) time from disease diagnosis to first biomarker testing was 266 (381) days, from the first biomarker testing to receipt of test results was 32 (101) days, and from biomarker testing to treatment initiation was 178 (394) days
 - Of all biomarker testing, 6.2% were single gene tests, 49.7% were broad panel tests, and 24.3% were single gene plus broad panel tests; for 19.8%, the type of test was unknown (**Figure 1**)
 - Among the patients with stage IV prostate cancer who received biomarker testing (n=1044), 265 (25.4%) tested positive for an actionable mutation and, of these, 35 (13.2%) received a targeted therapy (Figure 2)
 - For patients with stage I through IV prostate cancer who initiated treatment within the oncology network (n=6429), hormone therapy was the most common treatment received by patients with prostate cancer (n=5038, 78.4%), followed by chemotherapy, hormone therapy maintenance (n=635, 9.9%). Patients may have received supportive care as a part of these regimens (**Figure 3**)

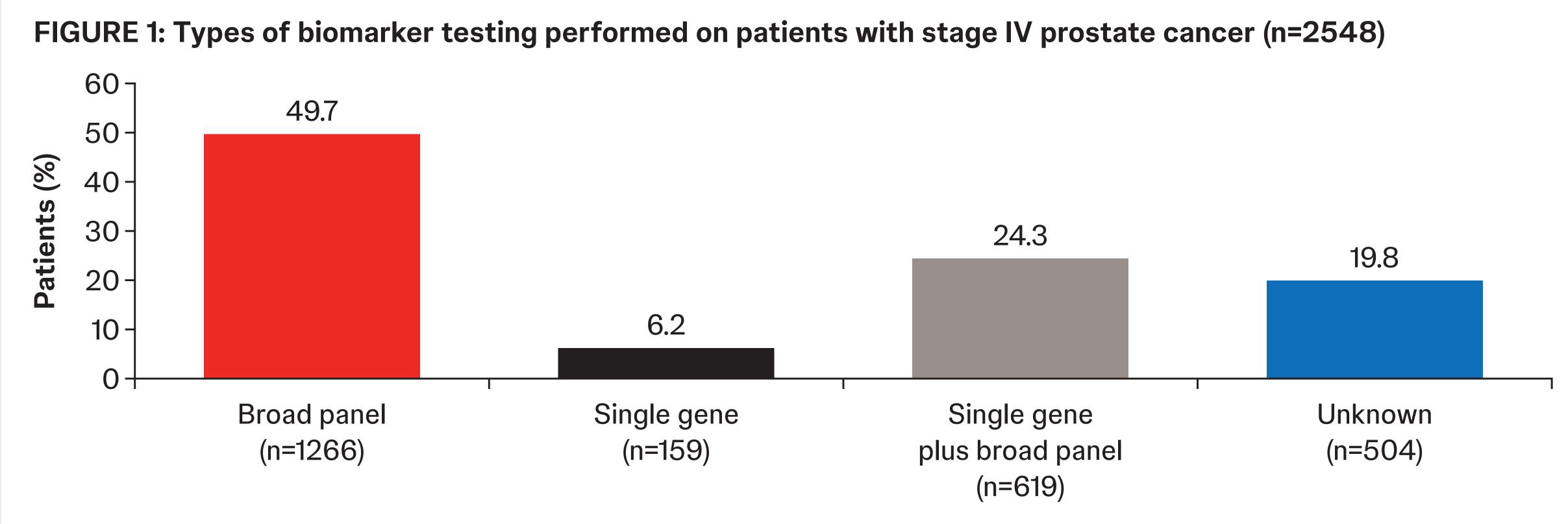
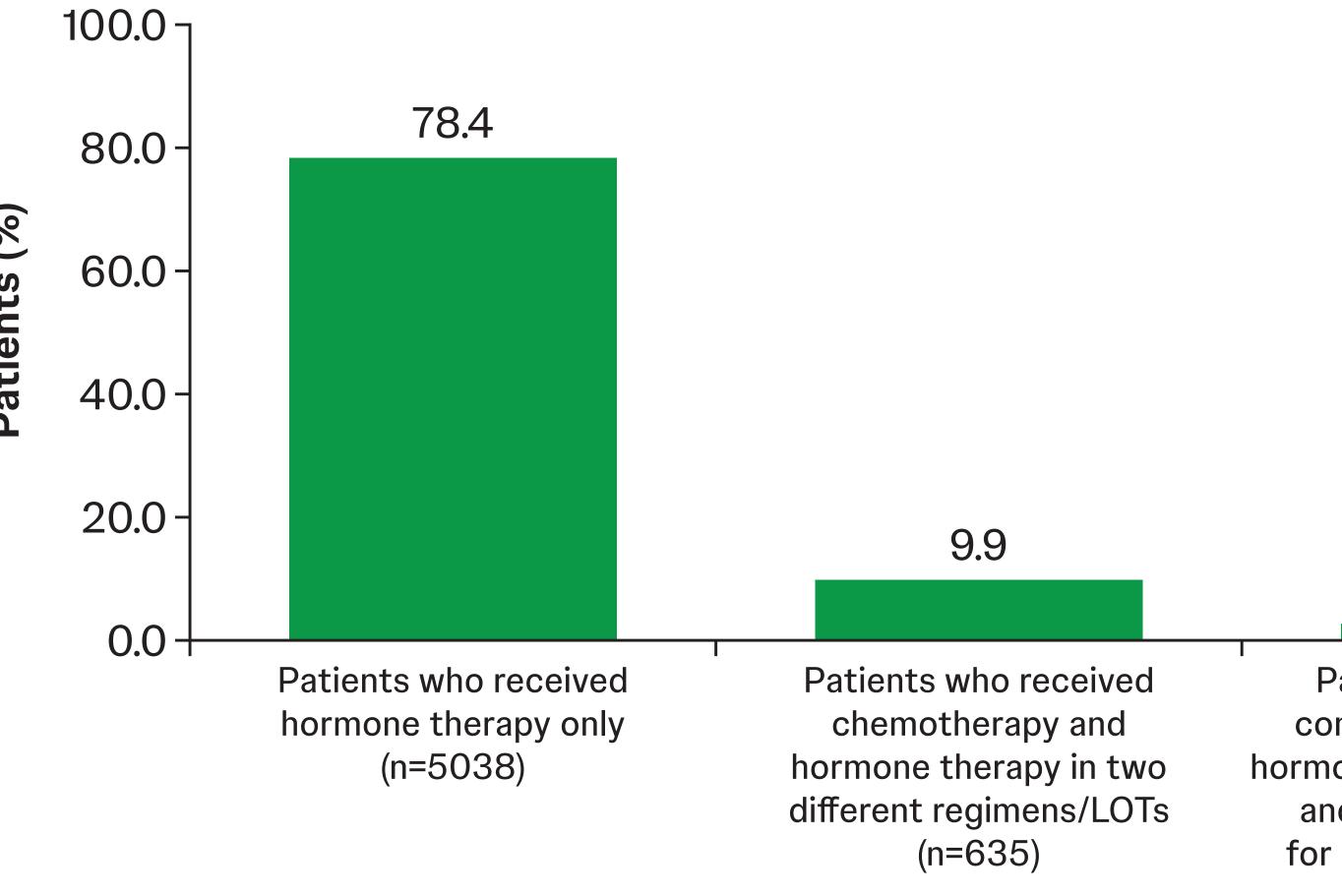


FIGURE 3: Treatments received by patients (n=6429)



database inputs consisted of iKnowMed (electronic medical records), ELLKAY CareEvolve (Genetic HL7 interface), and the oncology network's Molecular Data Warehouse, which covered

criptive statistics were used to report baseline demographics and clinical characteristics and to determine the percentage of patients with metastic disease who received biomarker testing

FIGURE 2: Numbers of patients with stage IV prostate cancer receiving biomarker testing, testing positive for an actionable mutation, and receiving targeted therapy (n=4510)

