

Utility of Oncotype Dx in Men with Intermediate-Risk Prostate Cancer

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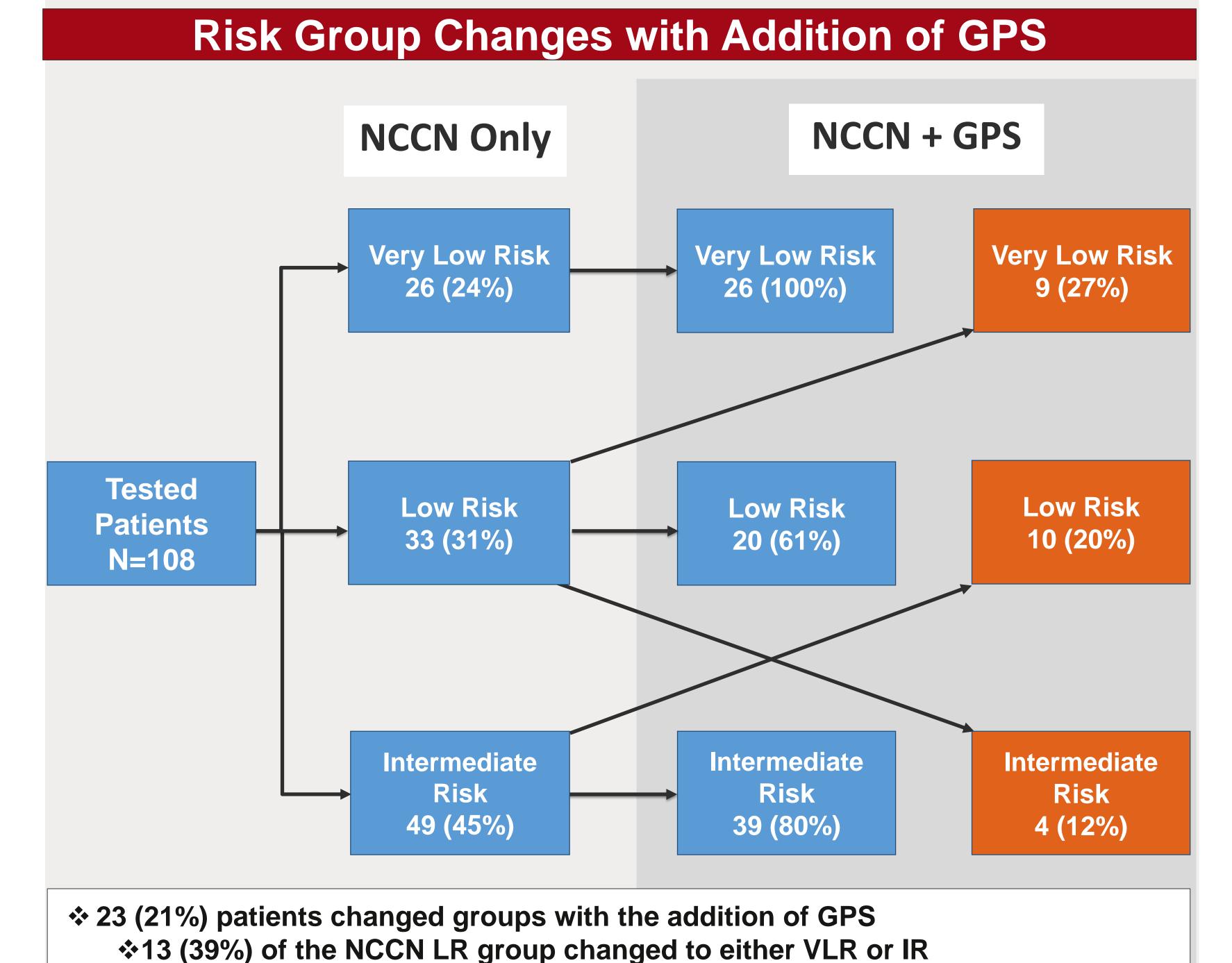
BACKGROUND

- ❖ The Oncotype Dx Genomic Prostate Score (GPS) Assay from Genomic Health [Redwood City, CA] is a commercially available assay designed for patients diagnosed with clinically localized prostate cancer to help guide treatment.
- Genomic Health has validated this test as an independent predictor of adverse pathology, and may predict PCSM and metastases following radical prostatectomy.
- ❖ Genomic Health recommends the test for men with lowand favorable intermediate risk prostate cancer [GS 3+3 and 3+4] who meet criteria for NCCN very low, low, or intermediate risk.
- ❖ Objective: To investigate the use of this test in clinical practice and assess its role in clinical management

METHODS

- ❖ Retrospective review of all Oncotype Dx reports received on prostate biopsies sent from a single institution to Genomic Health from 2015 to September 2017.
- Changes in patients' NCCN risk group [Very Low Risk (VLR), Low Risk (LR), or Intermediate Risk (IR)] as a result of the additional GPS were recorded.
- ❖ Decisions in disease managed were captured and included treatment by radial prostatectomy (RP), radiation therapy (XRT), active surveillance (AS), or focal therapy (FT).

RESULTS Prostate Biopsies Sent for Oncotype Dx 2015 — 2017 114 **Patients** (108 with sufficient tissue for testing) Age **50-83 years** Range Mean 65 years **Gleason Score** 64% (69/108) 3+3=6 36% (39/108) 3+4=7 **Number of Positive Cores** 1-18 Mean **Max Core Involvement** 76% (82/108) <50% 24% (26/108) ≥50% **Perineural Invasion** 13 (14/108)



❖10 (20%) of the NCCN IR group changed to LR

RESULTS

❖ 93 men had treatment information available

Interventions Selected Among Cohorts

NCCN + GPS	Patients	RP or XRT	AS	FT	Unknown	Comments
VLR	35	14% (5/35)	72% (25/35)	0	14% (5/35)	3 of 5 who received RP/XRT were LR by NCCN alone.
LR	30	10% (3/30)	70% (21/30)	3% (1/30)	17% (5/30)	10 men from changed from IR to LR, of which 8/10 chose AS.
IR	43	47% (20/43)	33% (14/43)	9% (4/43)	11% (5/43)	Of the 4 men who changed from LR to IR, 2 were definitively treated and 2 chose AS.

CONCLUSION

- ❖ Nearly half of the Oncotype Dx GPS assays sent at our institution were requested on patients who were intermediate risk by NCCN guidelines.
- The Oncotype Dx GPS assay did not change the risk group for the majority of men.
- Clinicians should discuss patient expectations and treatment preferences prior to performing additional genomic testing.
- Understanding how GPS may impact management decisions will improve the cost-effectiveness of such testing.