



OncoAssure™ Prostate

A Next Generation Risk Stratification Test for Assessment of Prostate Cancer Aggressiveness

The OncoAssure Prostate Test



Is used to **estimate the risk of having aggressive disease** (post-biopsy) or the **risk of biochemical recurrence** (post-surgery).



Employs a novel approach by utilizing master driver genes, which cover a wide range of biological pathways, to develop the prognostic signature; 4 prognostic (FOXM1, MCM3, MTUS1, TTC21B) and 2 reference (ALAS1, PPP2CA) genes.



Incorporates a well-known and well-validated clinical component; Cancer of the Prostate Risk Assessment (CAPRA), into the risk score.



Is prognostic for contemporary clinical features associated with tumor aggressiveness and poorer clinical outcomes, including the presence of cribriform morphology and Tertiary Grade 5.

Benefits of using OncoAssure Prostate

9.7 x
Risk of adverse pathology

Patients with a higher OncoAssure Prostate result score were 9.7 times more likely to experience adverse pathology to those with a lower OncoAssure Prostate result.



The risk of biochemical recurrence within 5 years was 28.2% for patients with a higher OncoAssure Prostate result, compared to just 6.4% for patients with a lower OncoAssure Prostate result.

Reduce unnecessary interventions >20%

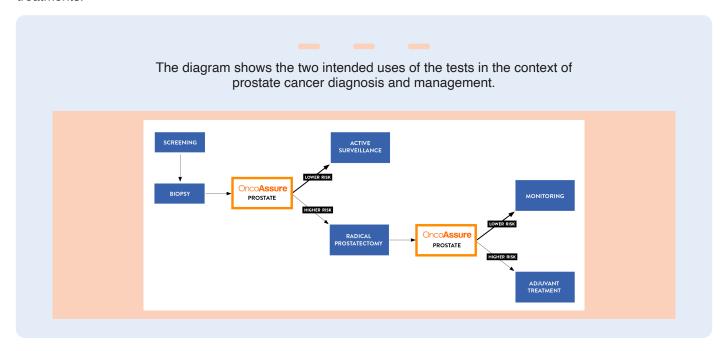
Using the OncoAssure Prostate test to guide treatment decisions instead of treating all patients, would lead to a 20% - 29% reduction in unnecessary interventions.

Introduction to OncoAssure Prostate Test

OncoAssure Prostate is a newly developed laboratory test (LDT) designed to help stratify risk in patients newly diagnosed with prostate cancer. It has demonstrated performance that is comparable to or better than existing market alternatives.

The OncoAssure Prostate test is a multigene prognostic signature incorporating clinicopathological information which assesses the probability of aggressive disease in men recently diagnosed with early-stage prostate cancer or men with localised disease who have undergone radical prostatectomy (RP).

Men whose OncoAssure Prostate risk score indicates they have a lower risk of aggressive disease may be suitable candidates for active surveillance. After RP, patients with a lower risk of biochemical recurrence can be reassured that surgery is sufficient to control their prostate cancer, whereas patients with a higher-risk of recurrence may need adjuvant treatments.



OncoAssure Prostate Test Development



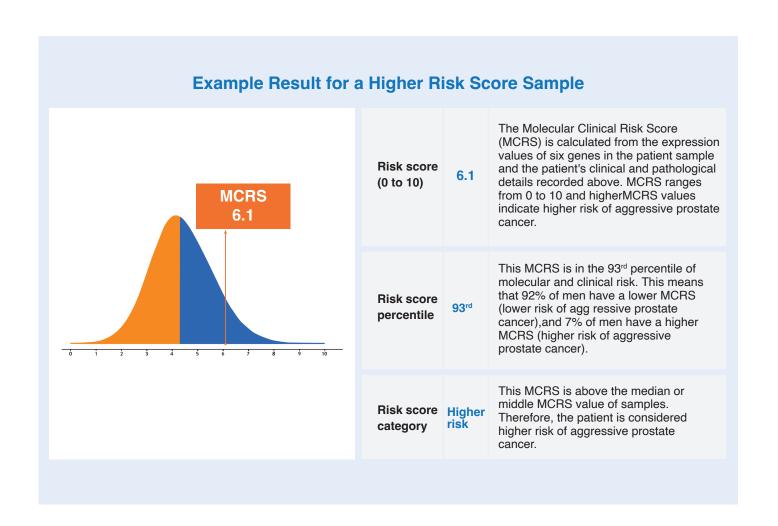
- 1: ESMO 2022 Paris, September 2022, Krzyzanowska et al. Annals of Oncol 2022, 33(S7):S1190
- 2: ASCO GU 2023 San Francisco, February 2023, Krzyzanowska et al. J Clin Oncol 2023, 41 (S6; Abstr 374)
- 3:Krzyzanowska A et al., Development, Validation, and Clinical Utility of a Six-gene Signature to Predict Aggressive Prostate Cancer. Eur Urol Focus. 2023. 9(6): p. 983-991
- 4: Krzyzanowska A, et al. Clinical validation of a biopsy-based six-gene signature prognostic for aggressive prostate cancer. BJUI Compass. 2025; 6(1):e474.



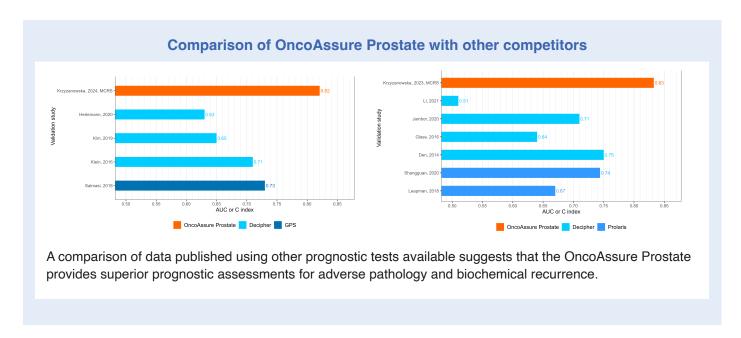
How to Use the OncoAssure Prostate Test

Based on the MCRS result on the report (see the interpretation example below), physicians can make recommendations on the treatment strategies (see the table below).

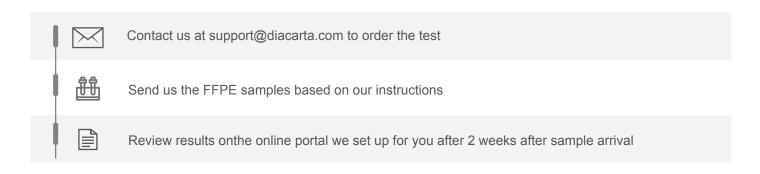
		NCCN Risk Category		
		Low/Very Low	Favorable Intermediate	Unfavorable Intermediate /High/Very High
(A) Onco Assure	Lower Risk Score	Consider active surveillance	Consider active surveillance	Consider treatment
	Higher Risk Score	Consider closer surveillance	Consider closer surveillance or treatment	Consider multimodal treatment



How Does OncoAssure Prostate Compare with 1st Generation Tests



Order OncoAssure Prostate



Insurance coverage: We build patient insurance or accept cash offers to cover the basic cost.

Test code: DC-19-1001 CPT code: 0497U

Disclaimer:

This laboratory developed test (LDT), OncoAssure™ Prostate, is performed at our CLIA-certified and CAP-accredited laboratory in Pleasanton, California. This test adds clinical value post-biopsy and post-surgery by providing prognostic information for adverse pathology and biochemical recurrence—key indicators of metastasis risk and long-term outcomes. This test has been analytically validated by the laboratory, but has not been cleared or approved by the U.S. Food and Drug Administration (FDA). For more information or to speak with a support specialist, please contact DiaCarta Clinical Laboratory at support@diacarta.com

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