Reduction in Therapeutic Burden from Use of CCP Test in Treatment Decisions among Newly Diagnosed Prostate Cancer Patients Independent of Charlson Comorbidity Index (CCI)

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OBJECTIVES

Risk of prostate cancer-specific disease progression and mortality may be assessed using the cell cycle progression (CCP) test.

RESULTS

The CCP score resulted in a change in Figure 1. Study Schema actual treatment administered in 48% of patients, with treatment changes stratified by CCI in Table 1. Physician Completes Part A-

- The Cell Cycle Progression (CCP) score was developed and validated to provide prognostic information to prostate cancer patients in all risk groups.^{1,2}
- A large prospective registry (P1000, n=1,206) recently showed that the CCP test results influence treatment decisions.³
- Here, we assess whether the CCI, an estimate of the severity of patient comorbid conditions, was correlated with these changes in treatment decisions.

METHODS

MOLECULAR TESTING

The CCP test is a validated molecular assay that

Physician IDs Eligible Patients

Initial Treatment Plan CCP Test Run on Patient Biopsy **CCP** Test Results Returned to Physician Physician Completes Part B-Intended Treatment

Physician Completes Part C-Agreed Upon Treatment

Physician Completes Part D-Actual Treatment

Our analysis shows that the CCP test result is a significant covariate, while the CCI is not (univariate: p=0.0153 vs p=0.1831, multivariate: p=0.0131 vs p=0.1528).

The CCI x Age interaction term is significant (p=0.0207), showing that CCI combined with age is correlated with changes in treatment.

However, the CCI x CCP score (p=0.5918) and CCI x Risk Percentile (p=0.5191) interaction terms are not significant.

- measures the RNA expression of 46 genes to generate a numeric CCP score.
- The CCP score is calculated by measuring the average RNA expression of 31 cell cycle progression genes normalized by the average expression of 15 housekeeping genes as quantified by RT-PCR.

STUDY DESCRIPTION

- P1000 included newly diagnosed (≤ 6 months) patients with untreated clinically localized prostate cancer.³
- Physicians completed a series of questionnaires to record their treatment recommendations (Figure 1), including:
 - Initial recommendations (pre-test)
 - Actual treatment (3–6 months of clinical follow-up)
- The CCI was recorded as part of the pre-test

Table 1. Treatment Change by CCI

CCI	Decrease	No Change	Increase	Total
0	289 (33.49%)	462 (53.53%)	112 (12.98%)	863
1	80 (37.74%)	105 (49.53%)	27 (12.74%)	212
2	24 (35.29%)	36 (52.94%)	8 (11.76%)	68
≥3	22 (34.92%)	33 (52.38%)	8 (12.70%)	63
Total	415	636	155	1206

CMH test (row-mean scores differ):

Unadjusted p=0.7933; Adjusted (risk percentiles): p=0.7546; Adjusted (AUA risk): p=0.9704

CONCLUSIONS

The lack of univariate and multivariate statistical significance implies that changes in treatment due to the CCP test are independent of CCI.



Changes in treatment decisions were examined with multinomial logistic regression modeling with CCI and CCP as continuous covariates along with other clinical









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