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Patient Reported vs. Claims Based Life Expectancy Tools: External Evaluation of Prediction Models in Men with Prostate Cancer

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Prostate Cancer

Figure 3. Leading Sites of New Cancer Cases and Deaths – 2020 Estimates

	Male			Female			
Estimated New Cases	Prostate	191,930	21%	Breast	276,480	30%	
	Lung & bronchus	116,300	13%	Lung & bron	chus 112,520	12%	
	Colon & rectum	78,300	9%	Colon & rect	tum 69,650	8%	
	Urinary bladder	62,100	7%	Uterine corp	bus 65,620	7%	
	Melanoma of the skin	60,190	7%	Thyroid	40,170	4%	
	Kidney & renal pelvis	45,520	5%	Melanoma d	of the skin 40,160	4%	
	Non-Hodgkin lymphoma	42,380	5%	Non-Hodgk	in lymphoma 34,860	4%	
	Oral cavity & pharynx	38,380	4%	Kidney & rei	nal pelvis 28,230	3%	
	Leukemia	35,470	4%	Pancreas	27,200	3%	
	Pancreas	30,400	3%	Leukemia	25,060	3%	
	All sites	893,660		All sites	912,930		
	Male						
	Male				Female		
	Male Lung & bronchus	72,500	23%	Lung & bron	Female chus 63,220	22%	
	Male Lung & bronchus Prostate	72,500 33,330	<u>23%</u> 10%	Lung & bron Breast	Female chus 63,220 42,170	22% 15%	
	Male Lung & bronchus Prostate Colon & rectum	72,500 33,330 28,630	23% 10% 9%	Lung & bron Breast Colon & rect	Female chus 63,220 42,170 um 24,570	22% 15% 9%	
aths	Male Lung & bronchus Prostate Colon & rectum Pancreas	72,500 33,330 28,630 24,640	23% 10% 9% 8%	Lung & bron Breast Colon & rect Pancreas	Female chus 63,220 42,170 um 24,570 22,410	22% 15% 9% 8%	
Deaths	Male Lung & bronchus Prostate Colon & rectum Pancreas Liver & intrahepatic bile duct	72,500 33,330 28,630 24,640 20,020	23% 10% 9% 8% 6%	Lung & bron Breast Colon & rect Pancreas Ovary	Female chus 63,220 42,170 um 24,570 22,410 13,940	22% 15% 9% 8% 5%	
ed Deaths	Male Lung & bronchus Prostate Colon & rectum Pancreas Liver & intrahepatic bile duct Leukemia	72,500 33,330 28,630 24,640 20,020 13,420	23% 10% 9% 8% 6% 4%	Lung & bron Breast Colon & rect Pancreas Ovary Uterine corp	Female chus 63,220 42,170 um 24,570 22,410 13,940 pus 12,590	22% 15% 9% 8% 5% 4%	
ated Deaths	Male Lung & bronchus Prostate Colon & rectum Pancreas Liver & intrahepatic bile duct Leukemia Esophagus	72,500 33,330 28,630 24,640 20,020 13,420 13,100	23% 10% 9% 8% 6% 4% 4%	Lung & bron Breast Colon & rect Pancreas Ovary Uterine corp Liver & intra	Female chus 63,220 42,170 um 24,570 22,410 13,940 pus 12,590 hepatic bile duct 10,140	22% 15% 9% 8% 5% 4% 4%	
timated Deaths	Male Lung & bronchus Prostate Colon & rectum Pancreas Liver & intrahepatic bile duct Leukemia Esophagus Urinary bladder	72,500 33,330 28,630 24,640 20,020 13,420 13,100 13,050	23% 10% 9% 8% 6% 4% 4% 4%	Lung & bron Breast Colon & rect Pancreas Ovary Uterine corp Liver & intra Leukemia	Female chus 63,220 42,170 um 24,570 22,410 13,940 pus 12,590 hepatic bile duct 10,140 9,680	22% 15% 9% 8% 5% 4% 4% 3%	
Estimated Deaths	Male Lung & bronchus Prostate Colon & rectum Pancreas Liver & intrahepatic bile duct Leukemia Esophagus Urinary bladder Non-Hodgkin lymphoma	72,500 33,330 28,630 24,640 20,020 13,420 13,100 13,050 11,460	23% 10% 9% 8% 6% 4% 4% 4% 4%	Lung & bron Breast Colon & rect Pancreas Ovary Uterine corp Liver & intra Leukemia Non-Hodgki	Female chus 63,220 42,170 um 24,570 22,410 13,940 pus 12,590 hepatic bile duct 10,140 9,680 8,480	22% 15% 9% 8% 5% 4% 4% 3% 3%	
Estimated Deaths	Male Lung & bronchus Prostate Colon & rectum Pancreas Liver & intrahepatic bile duct Leukemia Esophagus Urinary bladder Non-Hodgkin lymphoma Brain & other nervous system	72,500 33,330 28,630 24,640 20,020 13,420 13,100 13,050 11,460 10,190	23% 10% 9% 8% 6% 4% 4% 4% 4% 3%	Lung & bron Breast Colon & rect Pancreas Ovary Uterine corp Liver & intra Leukemia Non-Hodgki Brain & othe	Female chus 63,220 42,170 42,170 um 24,570 22,410 13,940 pus 12,590 hepatic bile duct 10,140 9,680 9,680 n lymphoma 8,480 er nervous system 7,830	22% 15% 9% 8% 5% 4% 4% 3% 3% 3%	

Estimates are rounded to the nearest 10, and cases exclude basal cell and squamous cell skin cancers and in situ carcinoma except urinary bladder. Estimates do not include Puerto Rico or other US territories. Ranking is based on modeled projections and may differ from the most recent observed data.

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What Do Prostate Cancer Patients Die Of?



2. Bill-Axelson A, Holmberg L, Garmo H, et al. Radical prostatectomy or watchful waiting in early prostate cancer. N Engl J Med. 2014;370(10):932–942.



Current Tools

- Comorbidity based Model (Cho) developed from Medicare claims
 - » Includes age, race, and Charlson Comorbidity Score
- Age Adjusted Comorbidity Model (Daskivich) developed in the VA population
 - » Includes age, race, treatment, PSA, Gleason Score, cancer stage, and the prostate cancer-specific comorbidity index
- Self Reported Health Status Nomogram (Hoffman): developed from Prostate Cancer Outcomes Study
 - » Includes age, race, and patient-reported overall health



Enthusiasm for Big Data

<u>Front Oncol</u> . 2016; 6: 149. Published online 2016 Jun 14. do	bi: <u>10.3389/fonc.2016.00149</u>	PMCID: PMC4905980 PMID: <u>27379211</u>					
Big Data Analytics for	Prostate Radiotherap	у					
James Coates, ^{1,*} Luis Souhami,	² and <u>Issam El Naqa</u> ³						
Author information > Article no	<u>J Am Med Inform Assoc</u> . 2015 No Published online 2015 Nov 9. doi	ov; 22(6): 1114. i: <u>10.1093/jamia/ocv136</u>	PMCID: PMC5009910 PMID: <u>26555016</u>				
The NIH Big Data to Knowledge (BD2K) initiative							
N C Med J. Author manuscript; Philip E Bourne, ^{1,*} Vivien Bonazzi, ¹ Michelle Dunn, ¹ Eric D Green, ² Mark Guyer, ¹ George Komatse Published in final edited form at Jennie Larkin, ¹ and Beth Russell ¹ N C Med J. 2014 Jul-Aug; 75 Author information > Copyright and License information Disclaimer							
Big Data for Populati	on-Based Cancer Res formation and Surveillance S	search System					
Anne-Marie Meyer, PhD, resear	ch assistant professor, Andrew F.	JAMA. 2018 Apr 3;319(13):1317-1318. doi: 10.1001/ja	ama.2017.18391.				
Laura Green, MBA, project manager, <u>Adrian Meyer</u> , MS, director PhD, MPH, assistant professor, Ethan Basch, MD, MSc, director,		Big Data and Machine Learning in Health Care.					
director	,, ,, ,, ,	Beam AL ¹ , <u>Kohane IS</u> ¹ .					
		Author information					
		PMID: 29532063 DOI: <u>10.1001/jama.2017.18391</u>					



Aim of the Study

To determine whether augmenting current life prediction tools with new data inputs (i.e. patientreported or claims-based health measures) improves predictive accuracy.



Methods

- Study Sample: SEER-CAHPS data
 - » Men 65 years and older diagnosed with prostate cancer from 2004 to 2013.
- Identified 3 existing tools that estimate life expectancy:
 - » Cho Model based on claims data
 - » Daskivich Model based on claims data
 - » Hoffman Model based on patient-reported health data
- Assessed incremental value of adding different data to predict other cause mortality using competing risk regression.



Results

- Among 3,240 men diagnosed with prostate cancer, 246 (7.62%) died of prostate cancer and 631 (19.48%) died of other causes.
- The 3 tools performed similarly well, with 10-year time-dependent AUCs ranging from 0.738 to 0.783.
- The addition of different data types modestly improved performance for all models.



Results

	Models	AIC	R ^{2*}	LR Test	DF	AUC
Cho model	Original model: Age, race and comorbidity level	9188.72	0.09	304.68	8	0.749
	Original model + Overall Health + Smoking	9115.30	0.11	386.09	12	0.753
Daskivich model	Original Model: Age, Race/Ethnicity, Primary Treatment, PSA, Gleason Score, Cancer Stage, PCCI category	9115.09	0.12	400.31	19	0.783
	Original Model + Overall Health + Smoking	9043.33	0.14	480.07	23	0.796
Hoffman model	Original Model: Age, Race/Ethnicity, Overall Health	9184.58	0.10	310.81	9	0.738
	Original Model + NCIComorbidity + Frailty	9117.34	0.11	388.06	14	0.744



Limitations

- Observational study
- Accuracy of coding practices
- Limited panel of patient-reported outcomes
- Unable to compare utility of different models due to limited number of patients with 10-years of follow-up
- Generalizability to non-Medicare populations



Conclusions

- In an external cohort, current life expectancy models performed similar to their initial description, serving as further evidence of their accuracy.
- Enriching models with new data types resulted in incremental improvements, suggesting that prediction can improve with increased data capture.
- Additional research will be needed to assess what combination of inputs provides the optimal balance of performance, feasibility, and meaningfulness.



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