

Prostate-specific antigen testing for prostate cancer screening: A national survey of Canadian primary care physicians' opinions and practices

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Cite as: *Can Urol Assoc J* 2017; Epub ahead of print. <http://dx.doi.org/10.5489/cuaj.4486>

Published online November 1, 2017

Abstract

Introduction: In 2014, the Canadian Task Force on Preventive Health Care (CTFPHC) recommended against routine prostate cancer screening with the prostate-specific antigen (PSA) blood test.¹ We surveyed Canadian primary care physicians (PCPs) to understand their opinions and attitudes towards prostate cancer screening in 2016.

Methods: Twenty PCPs piloted the survey to assess its accessibility. We distributed a flyer to 19 633 PCPs as an insert in a large mailed package inviting them to attend a national meeting, and later promoted the survey at the meeting. Multinomial logistic regression models examined factors associated with agreement of key guideline statements and the overall benefit of PSA screening.

Results: A total of 1254 PCPs responded (rate of 6.4%); 54.7% of physicians aware of the CTFPHC recommendations report screening less often as a result. Overall, 55.6% of PCPs feel that the risks of PSA screening outweigh the benefits. On multivariable analysis, physicians who did not read the guidelines, did not have an academic appointment, or were in practice for over 20 years were significantly more likely to disagree with the statement that men 55–69 years old should not be screened for prostate cancer with PSA.

Conclusions: Our national survey found that the prostate cancer screening practices of Canadian PCPs varies widely across physician demographic groups, with almost equal numbers for or against. This has significant ethical, medical, and legal implications. The poor response rate to highly incentivized survey request may suggest a reluctance or general apathy towards this subject because of the Task Force recommendations. Future efforts should provide physicians with objective guidance around PSA screening, incorporating input from all stakeholders, including PCPs, urologists, and patients.

Introduction

Prostate cancer is the most prevalent cancer amongst Canadian men, representing 24% of all new cancer diagnoses in Canada.¹ For nearly three decades, screening for prostate cancer with prostate-specific antigen (PSA) testing has been an essential component of preventive care.² More recently, the risks and benefits of PSA as a screening biomarker for prostate cancer have come under scrutiny, prompting a re-evaluation of its role in clinical practice.^{3,4}

First in 2008, and again in 2012, the United States Preventive Services Task Force (USPSTF)³ published recommendations against screening for prostate cancer based on two large randomized controlled trials. The USPSTF is a government-issued panel composed of clinical epidemiologists, internists and primary care physicians, who objectively analyze available data and make recommendations based on the perceived quality of the evidence. Following the 2012 updated recommendations, the Canadian government asked the Canadian Task Force on Preventive Health Care (CTFPHC)⁴ to undertake a comparable analysis. In 2014 a similar recommendation against screening for prostate cancer with the PSA test was published by the CTFPHC. The American and Canadian taskforces cite both of the large randomized trials in their recommendations, as neither was able to show an overall survival benefit in their screening arms despite evidence of false-positive biopsies, over-diagnosis of non-life threatening cancers, and subsequent complications from investigation and treatment.^{5,6}

These recommendations have been met with criticism from urologists, oncologists and patient advocacy groups. In October 2014, the Canadian Urological Association (CUA) issued a press release addressing the CTFPHC recommendations, citing concerns that the task force failed to include key observational studies that point to PSA's utility in both screening and risk-stratification of men aged less than 55 years old.^{7,8} The CUA also cited the failure to acknowledge the role of PSA screening in conservative management ("active surveillance") of diagnosed low risk cases,⁹ an established clinical practice in Canadian urology.

In light of this ongoing controversy, the views and practices of Canadian PCPs in 2016 remain heterogeneous. In 2012, Kapoor et al published the results of a provincial survey of Ontario family physicians, immediately following publication of the USPSTF guidelines.¹⁰ They found a wide variation amongst Ontario PCPs, around both general PSA screening practices and their individual beliefs about the utility of screening for prostate cancer. Earlier studies conducted in British Columbia and Newfoundland and Labrador found similar results.^{11,12}

We created a survey instrument to survey a national sample of Canadian PCPs to understand their knowledge of and agreement with the CTFPHC guidelines, their current screening practices, and their use of shared decision-making around PSA testing.

Methods

Our survey instrument was designed for distribution to PCPs across Canada who routinely see men of prostate cancer screening-appropriate age in their practice (Appendix-1). The survey questionnaire and content were developed systematically through multiple iterations, with input from experts in key stakeholder groups. Prior to distribution, a pilot survey was conducted using local area PCPs from both community and academic practices in Ontario and British Columbia, and feedback was collected regarding the content and accessibility of the instrument.

The population approached to complete the survey represented a sampling of PCPs from across Canada, excluding Quebec. A flyer was distributed as an insert to a mailed invitation to attend a national primary care physician conference. We used a raffle prize draw to incentivize participants to complete the survey. 19,633 physicians (of a total 30,902 PCPs in Canada, excluding Quebec) received the mailing including our invitation, informing them of the purpose of the study and directing them to a web address where the survey was located, hosted by Fluid Surveys (www.fluidsurveys.com). The survey was kept open for 3 months in total to ensure enough time for respondents to access the questionnaire (June-August 2016). We then re-opened and promoted the survey at the national conference, asking those who had yet to complete the questionnaire to do so.

Descriptive statistics included frequency-distribution data and histogram representation of survey responses. Stratification of the cohort allowed for data comparisons across different demographic categories. Mann-Whitney U and Kruskal-Wallis tests were used to identify significant variations in agreement with guideline statements and overall benefit of PSA screening across strata. We used multinomial logistic regression models to understand the relationship between physician demographic and practice type, agreement with guideline statements and interpretation of the overall risk-benefit relationship of PSA screening. Statistical significance was set at $p < 0.05$ based on a two-tailed comparison. Statistical analyses were performed using SAS 9.3 (SAS Institute Inc., Cary, NC, USA). Ethical approval was granted for the study (REB Ref. 670-1601-Uro-009).

Results

1,058 physicians accessed the survey after the mail-out flyer, with an additional 196 during the second enrollment period at the conference (total of 1,254 respondents; response rate 6.4%). 47 of these were excluded, as no responses were recorded. 17 respondents were excluded after reporting they were not currently working as PCPs in a Canadian practice or seeing men of screening age. A total of 1,190 responses were included in the final analysis (93% completion rate).

Demographics

Demographic data from both the first and second enrollment periods were similar, and so were combined for the overall analysis (Table-1).

Sources of information on screening guidance

Our questionnaire asked respondents to identify where they turn to for guidance regarding best practice in cancer screening. 45.4% reported using government agencies to inform their screening practice (e.g. CTFPHC), whereas 26.1% use specialist organizations (e.g. CUA), and 25.3% look to national or provincial Colleges.

Understanding of and agreement with CTFPHC guidelines

81.5% of respondents were aware of the 2014 CTFPHC guidelines at the time of the survey, 80.9% of whom reported having read the document. Of those reading the guideline, 78.1% perceived the guidelines to be either 'clear' or 'very clear'. Of those who were aware of the recommendations, 54.7% reported screening less as a result, 4.7% screen more often, and 40.5% reported no change in their screening practices (of whom 23.1% report not routinely using the PSA test).

We then asked respondents to state their level of agreement with the CTFPHC report's recommendations on screening for three separate age groups of men (Figure-1). There was little agreement regarding men aged 55 to 69 years of age, with 38.8% of respondents agreeing that men in this cohort should not be screened. Notably 10.6% of respondents reported disagreement with *all three* guideline statements, whereas 35% agreed with all CTFPHC recommendations.

Screening practice patterns

Respondents were asked to outline their screening practices for men with different prostate cancer risk profiles (Figure-2). We also inquired more generally about screening methods. 52.6% of respondents reported using both PSA and digital rectal examination (DRE), with 14.5% using DRE alone and 10.2% using only PSA testing without physical examination. Figure-3 illustrates the patient ages at which physician's initiate and terminate routine prostate cancer screening. When the initial test is normal the frequency of PSA testing by those who recommend screening was either annually (22.9%), every 2 years (31.6%) or not again (27.6%). Finally, we asked respondents to provide their overall level of agreement with the statement that in average risk men, PSA screening's benefits outweigh its risks (Figure-4).

Shared decision-making

The vast majority of respondents believe in a shared decision-making approach to PSA testing (Figure-4). 89.4% discuss the risks and benefits of screening with men, and 87.3% of physicians reported feeling comfortable having such a discussion with a patient in their practice. Respondents counsel their patients around many of the risks associated with PSA testing (Figure-5).

Multivariable analysis

Multinomial logistic regression models were constructed to better understand whether key demographics and practice-types of PCPs affected their agreement with guideline statements and overall perceived benefit of PSA screening (Table-2a). The relative odds of agreeing rather than being neutral was 2.07 (95% CI 1.39 to 3.09) times for PCPs with an academic appointment compared to PCPs without an academic appointment. Conversely, those with over 20 years in practice were more likely to disagree with this recommendation (OR 2.43, 95% CI 1.29-4.60). In men 55 to 69 years old, PCPs who had read the guidelines document were less likely to disagree than be neutral with the CTFPHC's recommendation (OR 0.59, 95% CI 0.37-0.95), as were those with an academic appointment (OR 0.67, 95% CI 0.46-0.98). However, physicians with 10-20 years experience were less likely to agree than be neutral with this recommendation (OR 0.58, 95% CI 0.36-0.96). Those with more than 20 years experience were also more likely to disagree than be neutral that men aged 55 to 69 should not be screened (OR 2.73, 95% CI 1.63-4.57). We examined which demographics predicted a PCP's agreement with the statement 'in average risk men (i.e. no risk factors for prostate cancer) the benefits of prostate cancer screening outweigh the risks' (Table-2b). Similar to the guideline statements, those PCPs who had read the guideline document were more likely to disagree than be neutral with this statement (OR 1.88, 95% CI 1.25-2.85). Physicians with greater than 20 years experience (OR 3.55, 95% CI 2.03-6.19) were more likely to agree than be neutral with the above statement, and those with 10-20 years experience were both more likely to agree (OR 2.00, 95% CI 1.12-3.58) and less likely to disagree (OR 0.59, 95% CI 0.36-0.95) that the benefits of PSA screening outweigh the risks, compared to physicians with neutral responses.

Discussion

Despite the disappointing response rate, our survey was able to collate the opinions and practices of over 1200 Canadian PCPs, and demonstrates the impact of the CTFPHC guidelines on prostate cancer screening in this country. As a result of simply being aware of the CTFPHC guideline, the majority (54.7%) of respondents state they have decreased the amount of screening they perform in their practice. As PCPs form the front line of cancer screening, this will have an undeniable impact on the number of men referred for biopsy and subsequently the incidence of prostate cancer diagnoses in Canada in future years. The vast majority of our respondents had no issue with the clarity of the guidelines document, with 78.1% stating they were 'clear' or 'very clear.' Additionally, we found that PCPs generally agreed with statements put forward by the CTFPHC, particularly that men under the age of 55 and over the age of 70 should not receive prostate cancer screening. However, it is worth noting that this general agreement does not hold true in the key demographic of men 55-69 years old.

Despite the general acceptance of these recommendations, there still exists a significant amount of variation amongst Canadian PCP's screening practices. This clearly has both medical and legal implications. Amongst average-risk patients, there is an even split in approach, with equal numbers of respondents recommending for and against PSA screening after a risk/benefit discussion with the patient. In addition, PCPs are screening men with lower urinary tract symptoms although these are not associated with an increased risk of prostate cancer diagnosis and this is clearly discussed in the CTFPHC recommendations. While it is difficult to answer why this is the case, this may represent a misunderstanding amongst physicians around current known prostate cancer risk factors.

The multinomial logistic regression models that were constructed allowed us to examine which respondent demographics and practice types predicted agreement with guideline statements and the overall benefit of PSA screening (Tables 2a & 2b). When looking at these analyses together, we can see that those PCPs with more years in practice seem to disagree with the CTFPHC's recommendations against PSA screening, and this same group of physicians also are more likely to agree with the notion that the benefit of PSA screening outweighs the risks overall. This finding is interesting as it is these physicians who were in practice before and during the initiation of PSA screening. It may be that the number of men presenting with locally advanced and metastatic prostate cancer encountered by this subgroup in the pre-screening era has dissuaded them from ceasing PSA screening despite CTFPHC recommendations.

Our survey results are compatible with recent observed trends in prostate cancer screening in both the United States and Canada. Bhindi et al. described a decrease in the number of men being referred for prostate biopsy to a high volume centre, in the wake of the 2012 USPTF recommendations¹³. They found that the detection rate of low-grade, but also intermediate and high-grade prostate cancers, dropped from 2008 to 2013 in their time series analysis. Similarly, studies from the United States also show that prostate cancer screening decreased following the 2008 USPTF recommendations,^{14,15} as did the incidence of low-grade prostate cancer diagnoses.^{16,17} These are expected epidemiological findings after the publication of a guideline against screening, but worryingly recent population data from the Surveillance, Epidemiology, and End Results (SEER) database indicates the rate of lethal cancers may be rising at the same time.¹⁸ Our study adds to this body of literature by addressing the perceptions and practices of PCPs, with whom the ultimate responsibility for carrying out PSA screening sits.

There are limitations to our survey, primarily the low response rate, and subsequently, possible non-response bias. Measures were taken to prevent this, such as piloting the survey medium, ensuring a long collection period (3 months), and using a generous incentive. Multiple studies have investigated the decline in physician survey response rates, citing survey burden and fatigue, perceived ineligibility, and lack of interest.¹⁹⁻²¹ Due to the nature of the survey distribution, we were unable to send reminder

notices, and this may have also contributed to the low response rate. We do not believe that the structure or content of the questionnaire itself contributed to this limitation, as the completion rate from those who accessed the survey was very high (93%). We are concerned that many PCP's are just not interested in the topic or do not deal with men during their at-risk years for prostate cancer. This apparent apathy should serve as a call to urologists, radiation and medical oncologists, and allied health care professionals to increase our efforts to engage PCP's in the prostate cancer screening conversation. Proposed avenues to accomplish this include the organization of community forums on the issue with urologist and PCP discussion and debate, academic and clinical collaborative efforts, and increased physician engagement with social media. A final limitation is that Quebec PCPs were not surveyed in this study because the flyers were not mailed to physicians in this province.

The gap between the Canadian Urological Association position and the perceptions of PCPs in our survey around prostate cancer screening appears to be wide. The results presented here show that there may be a disconnect in understanding between two groups of physicians crucial to the health of this population. Although the evidence for PSA screening is mixed, the rational and passionate arguments on both sides of the issue imply that the optimal, patient-centered approach to this problem lies somewhere between screening for all men or none. Our understanding of PSA has become more nuanced over the past two decades, and the evidence would suggest that careful patient selection and thoughtful timing of PSA testing (so-called "smart screening") can lead to fewer unnecessary biopsies and increased detection of high-risk cancers.²² While it is correct to look to high-level evidence for guidance on cancer screening, like the randomized control trials in this field, the unfortunate contamination of these studies mean that we must be cautious when interpreting their findings.²³ To make sweeping recommendations based on the results of these few studies at face value will see us return to a time when a diagnosis of prostate cancer often had a much bleaker presentation and outcome.²⁴

The landscape of prostate cancer screening continues to be in flux. A 2017 revision of the USPSTF recommendations saw this group's stance against PSA screening soften, changing their rating from a 'D' to a 'C' grade.²⁵ They cite evidence from long-term follow up in the ERSPC trial demonstrating improved cancer-specific survival (CSS),²⁶ and reduced metastatic disease burden²⁷ in the trial's screening arm. They also acknowledge the increasing acceptance of active surveillance in men with low-risk prostate cancer.²⁸ This change in recommendation immediately brought public and media attention to the issue,²⁹ and time will tell what impact this decision will have on PSA screening in Canada.

Conclusion

Our study demonstrates the impact the CTFPHC recommendations on prostate cancer screening have had on screening practices in Canada. More than half of Canadian PCPs reported being less willing to offer men screening with the PSA test. This has significant practice pattern, medical and legal implications, particularly if it results in a stage shift in diagnosis, with an increase of Canadian men presenting with metastatic disease. The low overall response rate of our survey must be considered when interpreting the responses. Despite this, the data presented here represents a diverse cohort of over 1200 Canadian PCPs. Our findings inform future work to monitor changes in prostate cancer care and emphasize the urgency for our urologic opinion leaders to provide all PCPs in Canada with clear, unified guidance.

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Figures and Tables

Fig. 1. Respondent agreement with routine screening in men of different age groups.

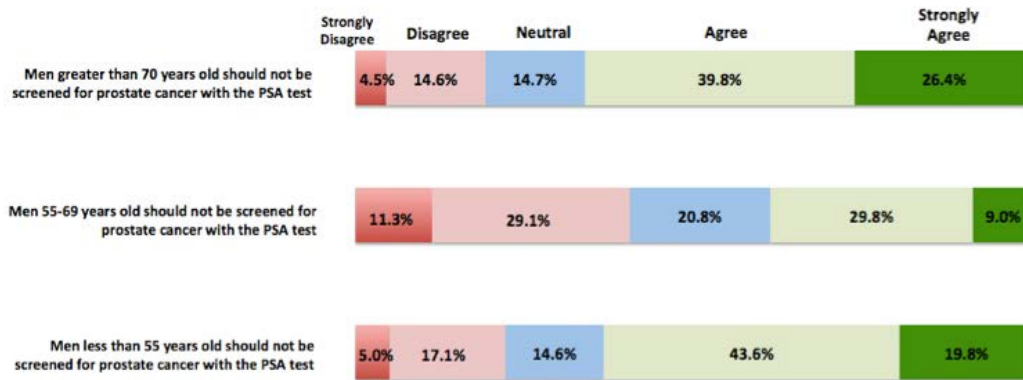


Fig. 2. Screening practices in men with different risk profiles.

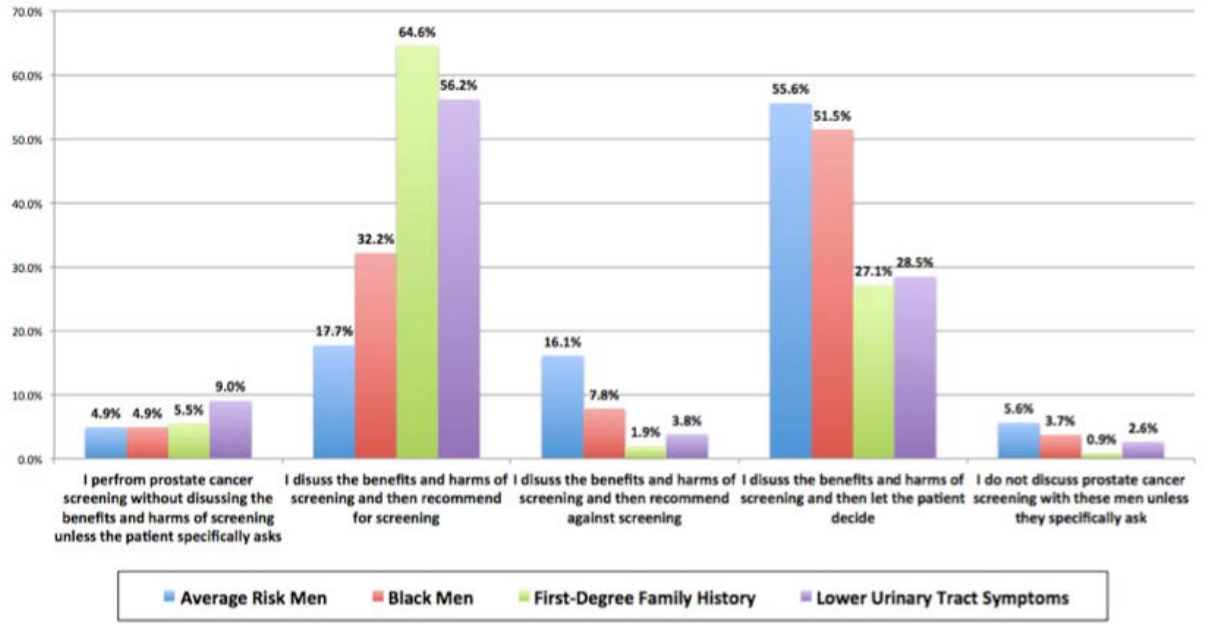


Fig. 3. At what ago do you start/stop offering routine cancer screening in average-risk men?

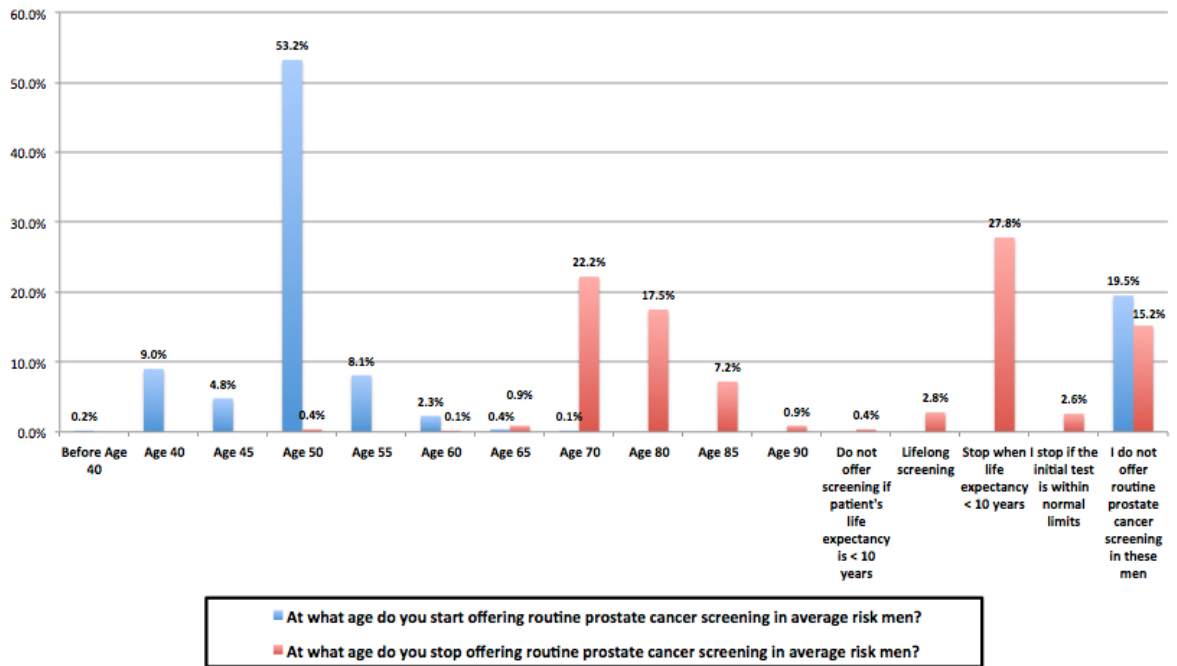


Fig. 4. Respondents' agreement with a shared-decision approach to screening.

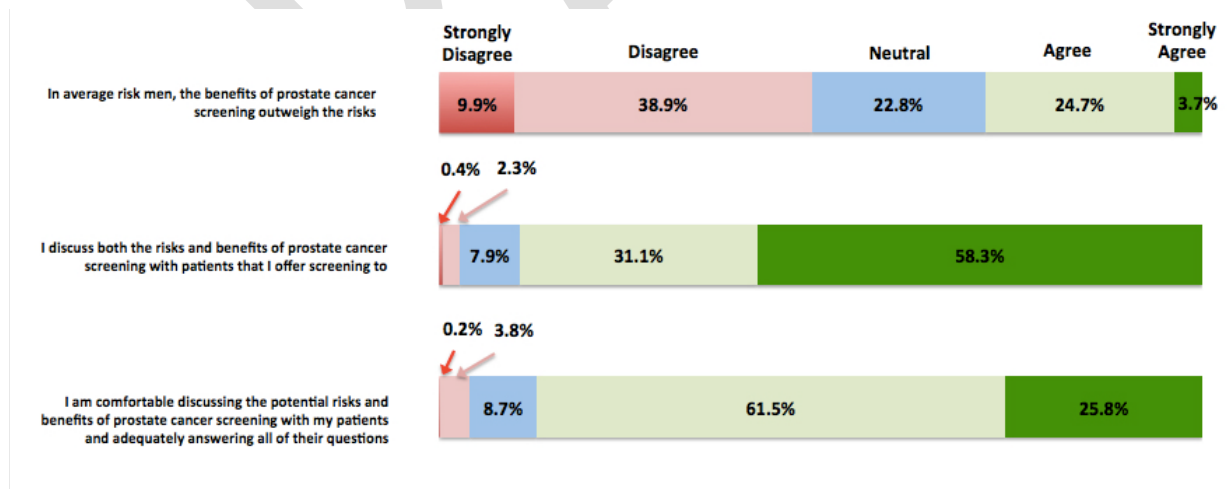
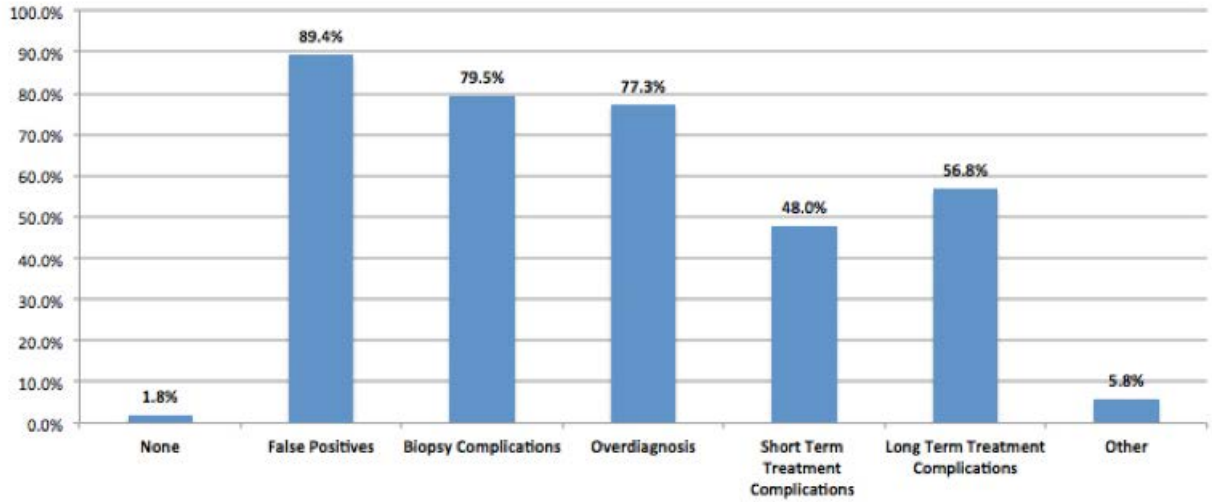


Fig. 5. Counselling patients on risks of prostate-specific antigen screening.

When discussing the PSA test and prostate cancer screening with patients, which potential harms of screening do you routinely mention?



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Table 1. Respondent demographics (n=1190)	
	No (%)
Gender	
Female	549 (46.1)
Male	641 (53.9)
Age	
<35 years old	332 (27.9)
35-44 years old	374 (31.4)
45-54 years old	218 (18.4)
55-64 years old	187 (15.7)
>65 years old	79 (6.6)
Province/Territory	
Alberta	216 (18.2)
British Columbia	309 (26.0)
Manitoba	40 (3.4)
New Brunswick	35 (2.9)
Newfoundland	18 (1.5)
Northwest Territories	1 (0.1)
Nova Scotia	31 (2.6)
Nunavut	2 (0.2)
Ontario	398 (37.3)
Prince Edward Island	5 (0.4)
Quebec	17 (1.4)
Saskatchewan	64 (5.4)
Yukon	5 (0.4)
Years in Practice	
<5	402 (33.8)
5-10	231 (19.4)
10-20	202 (17)
>20	355 (29.8)
Catchment area size	
Small population centre ($\leq 29,999$ people)	313 (26.3)
Medium population centre (30,000 – 99,999 people)	249 (20.9)
Large population centre ($\geq 100,000$ people)	628 (52.8)
Practice type	
Group practice	1020 (85.7)
Solo practice	170 (14.3)
Academic affiliation	
Yes	544 (45.7)
No	646 (54.3)

BC	1.17	0.47	2.95	0.86	0.39	1.91	1.01	0.49	2.05	1.02	0.50	2.08	0.77	0.26	2.29	0.41	0.16	1.05
Prairies	0.68	0.27	1.72	0.85	0.39	1.85	1.05	0.52	2.13	1.00	0.49	2.03	0.59	0.20	1.78	0.53	0.21	1.35
Central	0.63	0.26	1.56	1.03	0.48	2.20	0.67	0.34	1.34	1.26	0.64	2.48	0.46	0.15	1.37	0.71	0.28	1.76

Odds of agreeing or disagreeing with guidelines statements, compared to giving a neutral response. Bold values are statistically significant ($p < 0.05$). CI: confidence interval; OR: odds ratio; Ref: reference.

Table 2b. Results of multinomial logistic regression, part 2						
<i>Odds of agreeing or disagreeing that the benefits of PSA screening in average-risk men outweigh the risks, compared to a neutral response</i>						
	In average risk men (i.e., no risk factors for prostate cancer) the benefits of prostate cancer screening outweigh the risks					
	Disagree			Agree		
	OR	95% CI		OR	95% CI	
		Lower	Upper		Lower	Upper
Read guidelines (yes vs. no)	1.88	1.25	2.85	1.36	0.85	2.18
Academic appointment (yes vs. no)	1.51	1.08	2.13	0.83	0.56	1.24
Catchment area						
Small	Ref			Ref		
Medium	0.93	0.57	1.52	0.87	0.48	1.56
Large	0.87	0.58	1.30	0.94	0.58	1.52
Gender (female vs. male)	1.37	0.97	1.94	0.84	0.55	1.29
Years in practice						
< 5	Ref			Ref		
5 to 10	0.69	0.44	1.09	1.54	0.86	2.77
10 to 20	0.59	0.36	0.95	2.00	1.12	3.58
>20	1.04	0.65	1.65	3.55	2.03	6.19
Province						
Maritime + territories	Ref			Ref		
BC	0.99	0.50	1.96	0.93	0.44	1.96
Prairies	1.06	0.54	2.08	0.69	0.32	1.45
Central	1.40	0.73	2.70	0.55	0.26	1.14

Bold values are statistically significant ($p < 0.05$). CI; confidence interval; OR: odds ratio; Ref: reference.