Shared decision making for men facing prostate cancer treatment: a systematic review of randomized controlled trials

Aims: To synthesize the empirical evidence on the effectiveness of shared decision making (SDM) compared to usual care for prostate cancer (PC) treatment.

Methods and results: A systematic review of academic (MEDLINE, EMBASE, Cochrane Library, CINHAL, PsychINFO, and Scopus) and grey (clinicaltrials.gov, WHO trial search, meta-Register ISRCTN, Google Scholar, opengrey, and ohri.ca) literature, also identified from contacting authors and hand-searching bibliographies. We included randomized controlled trials (RCTs): 1) comparing SDM to usual care for decisions about PC treatment, 2) conducted in primary or specialized care, 3) fulfilling the key SDM features, and 4) reporting quantitative outcome data. Four RCTs from Canada (n=3) and the USA were included and comprised 1,065 randomized men, most (89.8%) of whom were in PC stage T1-T2. The studies reported 24 outcome measures. In 62.5% study estimates, SDM was similar to usual care at improving patient satisfaction and mood, and at reducing decisional conflict and decisional regret. In 37.5% study estimates, SDM significantly improved knowledge, perception of being informed and patient-perceived quality of life (QoL) at four weeks. There was a dearth of outcome data, particularly on the adherence to treatment and on patient-important and clinically relevant health outcomes such as symptoms and mortality.

Conclusion: SDM may positively influence men’s knowledge and may have a positive but short-term effect on patient-perceived QoL. The (long-term) effects of SDM on patient-related outcomes for decisions about PC treatment are unclear. Future research needs consensus about the interventions and outcomes needed to evaluate SDM and should address the absence of evidence on health outcomes.

Keywords: systematic review, shared decision making, prostate cancer, treatment, controlled clinical trials, urology

Introduction

Prostate cancer (PC) is the second leading cancer in men and the fifth leading cause of death due to cancer in men worldwide. Patients with PC often face more than one alternative to treatment eg, active treatment, active surveillance, or watchful waiting. These choices involve trade-offs between benefits and harms due to the limited evidence regarding the optimal treatment strategy for PC. The survival benefit of treatment options including surveillance is associated with considerable morbidity due to potential adverse outcomes of treatment (eg, urinary and erectile dysfunction, loss of fertility, and chemotherapy and/or hormone therapy side
These factors make treatment decisions complex and highly preference-sensitive. Patients thus need to weigh carefully not only the diagnoses and prognoses but also their own fears, values, beliefs, ethics, hopes, and previous experience.

Shared decision making (SDM) is viewed as an approach to involve patients and their clinicians in a process of collaboration and deliberation to reach medical decisions, particularly for preference-sensitive conditions. SDM helps inform patients about the options for, and the effectiveness of, treatment, taking into account the patient’s needs, knowledge and their value of risks, benefits, and harms. Health authorities and policy makers strongly encourage SDM for decisions about PC treatment. There is variation in the level of SDM implementation however, mainly due to the lack of consensus in SDM definition and goals. We performed a systematic review to evaluate the evidence for the effectiveness of SDM compared to current clinical practice for the treatment of PC.

**Methods**

We followed a protocol based on the principles for systematic reviews and report the methods according to the PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) guidelines (see Table S1). We identified and selected studies following the methods published in detail elsewhere.

**Eligibility criteria**

We included peer-reviewed and grey literature from RCTs published in English comparing the effects of SDM to usual care for decisions about PC treatment, which were conducted in primary and specialized care (general practice, community clinics, ambulatory care, hospital and private). We further limited the inclusion of studies to RCTs that, regardless of the intervention target (patients, HCP, surrogates, or family members): 1) met the criteria for SDM, supporting the principle of bi-directional deliberation, as previously illustrated, and 2) reported data in quantitative format for the outcomes of knowledge, patient satisfaction, perception of being informed, risk perception, decisional conflict, decisional regret, outcomes of emotional distress caused by the disease itself and/or treatment (eg, mood disturbance, anxiety, and depression), treatment behavior (eg, adherence to treatment), quality of life (QoL), symptoms, and mortality.

**Study identification and search strategy**

We searched for studies in: 1) academic databases: MEDLINE (Ovid), EMBASE (Elsevier), CINHAL (EBSCOHost), The Cochrane Library (Wiley), PsychINFO (EBSCOHost) and Scopus up to March 2015 (see Table S2); and 2) grey literature records (accessed: Feb-Aug, 2016) from clinicaltrials.gov and ISRCTN, the WHO search portal (http://apps.who.int/trialsearch), the Ottawa Hospital Research Institute website (http://www.ohri.ca), Google Scholar, and the system for Information on Grey Literature in Europe (http://opengrey.eu/). We additionally searched for the registration numbers of the trials using Medline and PubMed. We screened the reference lists of included studies, relevant reviews, and clinical guidelines, and contacted (Jun 2015–Jan 2017) the authors of abstracts for which the full-texts could not be located.

**Selection of studies and data extraction**

Two independent reviewers screened all titles and abstracts, and assessed the full-text of eligible publications. One reviewer extracted all data and a second reviewer independently verified data extractions. Differences in study selection and data extractions were resolved by consensus or by involving an arbitrator. We grouped outcomes into affective-cognitive, behavioral, and health outcomes following the system by Shay and Lafata (2015).

**Risk of bias**

Two reviewers independently assessed in duplicate the quality features of the included studies following established guidelines and resolving differences by discussion. We rated the adequacy of core items including generation of random sequence, concealment of allocation at randomization, blinding (patients, health care providers, and outcome assessors), intention-to-treat (ITT) (if participants were analyzed based on their original group allocation), follow-up (same length of time), and if there were attrition rates of significant concern (at least 20%). We also rated whether studies reported on the definition of inclusion and exclusion criteria, “a-priori” sample size calculation, primary and secondary outcomes, and funding sources.
**Data synthesis**

There was mostly one study per outcome precluding the ability to perform meta-analyses. We thus analyzed the data based on individual trial estimates. Where data were sufficiently reported, we calculated the unadjusted risk ratios (RR) or the standard mean differences (SMD) and the 95% confidence intervals (CI) assuming a random-effects model in RevMan 5.3.5 software.  

**Results**

We examined in detail the full-text of 270 articles. Four of these fulfilled the SDM criteria and investigated the comparative effectiveness of SDM with usual care (Figure 1).

**Study and population characteristics**

The studies were from Canada (n=3) and the USA (n=1), and comprised 1,065 patients individually randomized to the
intervention groups (Table 1). The patients were recruited from multidisciplinary or specialized care in hospital or general medicine. Three RCTs reported on cancer staging: 89.8% of the men had localized PC (clinical stage T1-T2),22–24 6.2% were in advanced stage (T3-T4),23 and 4.1% were of “unknown” clinical stage.22,23 Radical prostatectomy (36.7%) and watchful waiting (28.5%) were the most frequently selected treatments among 743 men from 2 RCTs, 1 in general (hospital) and 1 in specialized cancer care, respectively. “No treatment” accounted for 25.4% of all options offered in one study.23 Men had a mean age of 64.6 (SD 7.8), 83.9% were married and at least 53.4% had a minimum of high school education. The only RCT reporting on ethnicity included White (71.5%) and Black (28.5%) men. In three RCTs reporting on employment status, 55.3% of the men were in full- or part-time employment. In two RCTs, 88.2% of the participating health care providers were radiation oncologists and 11.8% were urologists.

Interventions’ characteristics
All interventions fulfilled the key features of SDM as illustrated in detail elsewhere.14 All RCTs used patient-directed interventions; one included the patients’ partners in the interventions sessions (Table 1).25 The interventions were delivered before decision making, within the time of scheduled visits23 or before consultations22,24,25 in order to empower patient participation in decisions. Men were recommended to review the material before consultation or were specifically encouraged to participate in treatment decisions. Men were advised to bring their significant other(s) to the consultation in one RCT.25 Three RCTs used multifaceted interventions on-site22,25 or at home.24 The interventions were self-administered or delivered by the research staff or by a nurse. The formats of the interventions included video, printed paper-based material, interviews, telephone calls, or audiotape recording. Only one RCT considered health care literacy for the development or pilot testing of the interventions.24 One RCT evaluated two SDM interventions and usual care.24 The content of the interventions included educational information, eg, about PC, treatment choices, advantages and disadvantages of treatment, side effects, and prognosis.

Risk of bias in the methods of included studies
All trials adequately randomized patients, but only two reported adequate concealment of allocation, leading to risk of selection bias in the other trials (Table 2). Only one RCT blinded patients and physicians, another blinded patients only, and no study performed or reported on the blinding of outcome assessors, resulting in high risk of performance and detection biases. Two RCTs with an attrition rate of less than 20% did not report on ITT analyses. Two RCTs had unclear reporting of attrition rates and ITT techniques. Men in all intervention arms were followed-up for the same length of time. All trials reported the participants’ inclusion criteria but only one trial reported the exclusion criteria too. Two RCTs reported on the sample size calculation and power. All RCTs measured the success of interventions by definition of primary outcome(s). All RCTs were funded by non-profit organizations.

Effectiveness of interventions
There were twenty-four outcome measures reported in the four RCTs (Figure 2). Data were sparsely reported with one study per outcome, limiting the ability to conduct meta-analyses. Table 3 shows the effect estimates for each individual trial.

Affective-cognitive outcomes
Knowledge
One trial implemented two SDM interventions.24 The different components between SDM groups consisted of nurse telephone calls to patients (treatment direct [TD]) or nurse telephone calls to patients and primary supporting persons (treatment supplement [TS]). Compared to usual care, TD significantly improved knowledge for PC and treatment at four weeks (SMD 0.33, 95% CI 0.03 to 0.64, p=0.03) and at three months (SMD 0.35, 95% CI 0.04 to 0.66, p=0.02). TS also showed a small but significant improvement in knowledge at four weeks (SMD 0.33, 95% CI 0.02 to 0.64, p=0.04), but not at three months. The combined effect of both SDM (TD and TS) interventions showed a significant improvement in knowledge at four weeks (SMD 0.35, 95% CI 0.08 to 0.62, p=0.01) and at three months (SMD 0.31, 95% CI 0.04 to 0.58, p=0.02).

Patient satisfaction
Individual trial effect estimates showed no significant difference between SDM and usual care in the number of patients who were satisfied with their treatment choice or who were satisfied with the levels of involvement in treatment decision making with the doctor.22 Scores of patient
### Table 1 Characteristics of included studies

| Author, year, country | Health care context, setting and facilities, n | Target population | Interventions (s) & randomized patients, N | Comparator(s) & randomized patients, N | Age: mean (SD) & target (range), years | Race or ethnicity, % | Married, % | Education, % | Employed, % | HCP |
|-----------------------|-----------------------------------------------|-------------------|--------------------------------------------|------------------------------------------|----------------------------------------|---------------------|------------|--------------|-------------|-----|
| Mishel et al, 2009†‡‡  | Multidisciplinary (hospital and specialized) Cancer center, n=2 Community hospital, n=3 Veterans’ medical center, n=1 | Men with biopsy-confirmed localized PC, no major cognitive impairment, ability to read, access to telephone, no prior cancer history, and a patient-designated PSP willing to participate in the study | NR | 1) Treatment supplemented: Communication strategy DVD + Information Booklet + 4 Telephone calls by same nurse to patients and PSP; N=89 2) Treatment direct: Communication strategy DVD + Information Booklet + 4 Telephone calls by nurse; N=93 | 62.5 (7.4) | 71.5 | 28.5 | 0 | 80.8 | 15.1 (SD 3.4) years of education | 56.6 | NR |
| Hack et al, 2007† | Specialized (Cancer) Tertiary oncology clinic treatment facilities, n=4 (in 3 cities) | Men with confirmed diagnosis of PC, presenting to a tertiary oncology clinic for their primary treatment consultation, without cognitive impairment disabling them from providing informed consent | • Radical prostatectomy (10.7)  • Hormone therapy (18.0)  • Watchful waiting (43.9)  • No treatment (25.4)  • Other (2.0) | No audiotape: 1) Consultation not audiotaped: no audio recording of primary treatment consultation, N=13; and 2) Not given audiotape: audio recording without giving audiotape, N=98 | 67.0 (8.0) (>18) | NR | NR | NR | 84 | 43.5% with more than HS | NR | Radiation oncologists, n=15 |

(Continued)
| Author, year, country | Health care context, setting and facilities, n | Target population | Patients with chosen treatment & randomized patients, N | Intervention(s) & randomized patients, N | Comparator(s) & randomized patients, N | Age: mean (SD) & target (range), years | Race or ethnicity, % | Married, % | Education | Employed, % | HCP |
|-----------------------|--------------------------------------------|------------------|---------------------------------|---------------------------------|---------------------------------|---------------------------------|----------------|-------------|-------|-----------|-----|
| Davison et al, 200722 | Hospital care General hospital, n=1 | Men with biopsy-confirmed localized PC, aware of their diagnosis, who had an initial urologic treatment consultation, but not scheduled for definitive treatment within the following 4 weeks, and able to read and write English | Men's partners were included in the sessions if they accompanied the patient | Generic information 24-min videotape with section on the identification of men's preferences + Information printout individualized according to patient's disease characteristics + Written information package with list of relevant info and help sources; N=162 | Men encouraged to bring PSP and were included in the sessions if present | 62.4 (6.9) | NR | NR | 85.4 | 68.8% with more than HS | 59.3 | NR |

(Continued)
| Author, year, country | Health care context, setting and facilities, n | Target population | Patients with chosen treatment † (%) | Intervention(s) & randomized patients, N | Comparator(s) & randomized patients, N | Age: mean (SD) & target (range), years | Race or ethnicity, % | Married, % | Education | Employed, % | HCP |
|-----------------------|-----------------------------------------------|-------------------|-------------------------------------|----------------------------------------|----------------------------------------|------------------------------------------|------------------|--------------|------------|-------------|------|
| Davison et al, 1997 | General medicine Community clinic with practicing urologists, n=1 | Men newly diagnosed with PC (stage not specified), aware of their diagnosis, who had not have an initial treatment consultation, with no evidence of mental confusion, able to read, speak and write English | Pelvic lymph node dissection (NR) | Written information package + Blank audiotape (optional) to record consultation + Interview with nurse with elements of patient empowerment to prepare men for consultation and to encourage participation in DM; N=30 | Written information package only; N=30 | 67.9 (7.1) (41–81) | NR | NR | 86.6 | 41.7% with at least HS Overall: 10.8 (SD 2.8) years of education | 28.3 | Urology physicians, n=2 |

Notes: † Hack, 2007: 449 patients received/selected treatment by the time of consultation; “Other” = transurethral resection of the prostate, orchiectomy. Davison, 2007: 294 patients. Davison, 1997: 75% of the men started/received treatment by the time of second interview.

Abbreviations: DM, Decision Making; HCP, Health care Professional(s); HS, High School; NR, Not Reported; PC, Prostate Cancer; PSP, Primary Support Person (e.g., wife).
Table 2 Methodological features of included studies

| Author, year | Country & funding | Outcome definition | Participants criteria | Study size | Sample size calculation and power | Groups comparable at baseline | Adequate sequence generation | Adequate allocation concealment | Adequate blinding | Attrition (outcome), % (range) | Adequate follow-up | ITT data |
|--------------|-------------------|--------------------|----------------------|------------|-----------------------------------|-----------------------------|-----------------------------|-------------------------------|-----------------|--------------------------------|----------------|----------|
| Mishel et al, 2009<sup>24</sup> | USA, non-profit  | Primary & Secondary | Inclusion: yes Exclusion: yes | N>200 nr | partially<sup>1</sup> | Yes | Unclear | HCP: unclear Participants: unclear OA: unclear | Unclear | Yes | Unclear |
| Hack et al, 2007<sup>23</sup> | Canada, non-profit | Primary | Inclusion: yes Exclusion: nr | N>200 | Yes | Yes | Yes | Yes | HCP: yes Participants: yes OA: nr | <20% | Yes | No |
| Davison et al, 2007<sup>22</sup> | Canada, non-profit | Primary | Inclusion: yes Exclusion: nr | N>200 | Yes | Yes | Yes | Yes | HCP: unclear Participants: yes OA: nr | <20% (9.26-9.57) or unclear | Yes | No |
| Davison et al, 1997<sup>21</sup> | Canada, non-profit | Primary | Inclusion: yes Exclusion: nr | N<200 nr | Yes | Yes | Unclear | HCP: no Participants: no OA: no | Unclear | Yes | Unclear |

Notes: <sup>1</sup>Patients in the control group had higher level of education than patients in the treatment direct group.

Abbreviations: HCP, Health care Professionals; OA, Outcome Assessors; nr, not reported.
satisfaction with preparation for decision making were not significant between groups.

**Perception of being informed and risk perception**

Individual trial effect estimates showed a small but significant effect of SDM when compared to usual care on patients’ perception of receiving information (SMD 0.20, 95% CI 0.01 to 0.39, p=0.04). The same trial showed a marginal but statistically significant effect of SDM on positive perceptions of receiving information about treatment alternatives (SMD 0.19, 95% CI –0.00 to 0.38, p=0.05) and treatment side effects (SMD 0.25, 95% CI 0.06 to 0.44, p=0.010). No study reported on risk perception.

**Mood state/disturbance, anxiety, and depression**

Outcomes of emotional distress including mood/disturbance, anxiety, and depression caused by the personal situation, the disease itself, or treatment were scarcely reported. Individual trial effect estimates showed no significant differences between SDM (TD or TS) and usual care with respect to mood state or disturbance at four weeks or at three months. No study reported quantitative data on anxiety and depression.

**Decisional conflict**

Individual trial effect estimates showed no significant differences between SDM and usual care in the scores of decisional conflict.

**Decisional regret**

Individual trial effect estimates showed no significant differences between SDM (TD or TS) and usual care in the scores of decisional regret at three months. The combined effect of both SDM (TD and TS) interventions compared to usual care was not significant at three months.

**Behavioral outcomes**

**Adherence to treatment of choice**

Radical prostatectomy and watchful waiting were the most frequent treatments received or chosen by men in two RCTs. No trial reported on the adherence to a treatment initially chosen however.

**Health outcomes**

**Quality of life, symptoms, and mortality**

Individual trial effect estimates showed a significant effect of two SDM (TD or TS) interventions, compared to usual care, on men’s positive feelings about their QoL at four weeks (TD: SMD 0.50, 95% CI 0.19 to 0.81, p=0.002; TS: SMD 0.50, 95% CI 0.18 to 0.81, p=0.002). This effect did not sustain at three months for either of the SDM (TD or TS) interventions however. Health outcomes of symptoms and mortality were not reported.

**Discussion**

In this systematic review, we identified four RCTs that both fulfilled the criteria for SDM and evaluated the comparative effectiveness of SDM with usual care for men faced with decisions about PC treatment.

Despite the low volume of literature, 37.5% of the study estimates significantly favored SDM when compared to usual care. We found moderate effects of SDM on knowledge, perception of being informed, and QoL. The effects on knowledge and QoL did not sustain at long term (three months) however. The remaining 62.5% of the study...
### Table 3 Outcome effect estimates for SDM compared to usual care for decisions about prostate cancer treatment

| Author, year | Outcome | Measurement method/tool | Measurement point | Intervention | Control | Effect estimate |
|--------------|---------|-------------------------|-------------------|--------------|---------|-----------------|
| **BINARY DATA** |         |                         |                   | SDM | Patients, n | Total patients, N | Usual Care | Patients, n | Total patients, N | RR (95% CI) |
| Davison et al, 2007 | Satisfaction with treatment choice | 10-item questionnaire based on 5-point Likert scale | 4–10 weeks after visit - after men had made a treatment decision with/out their urologist | Generic information 24-min videotape with section on the identification of men’s preferences + Information printout individualized according to patient’s disease characteristics + Written information package with list of relevant info and help sources + Telephone calls by RN approx. 4 weeks after visits | 141 | 149 | Generic information 24-min videotape + Written information package with list of relevant info and help sources | 135 | 145 | 1.02 (0.96 to 1.08) |
| Davison et al, 2007 | Satisfaction with level of involvement in decision making | 5-item questionnaire based on 5-point Likert scale | 4–10 weeks after visit - after men had made a treatment decision with/out their urologist | Generic information 24-min videotape with section on the identification of men’s preferences + Information printout individualized according to patient’s disease characteristics + Written information package with list of relevant info and help sources + Telephone calls by RN approx. 4 weeks after visits | 142 | 149 | Generic information 24-min videotape + Written information package with list of relevant info and help sources | 133 | 145 | 1.04 (0.98 to 1.10) |

(Continued)
Table 3 (Continued).

| Author, year | Outcome | Measurement method/tool | Measurement point | Intervention | Control | Effect estimate |
|--------------|---------|-------------------------|-------------------|--------------|---------|-----------------|

**CONTINUOUS DATA**

|                     |        |                         |                   |              |         |                 |
|---------------------|--------|-------------------------|-------------------|--------------|---------|-----------------|
| **SDM**             | mean (SD) | Total patients, N | Usual Care | mean (SD) | Total patients, N | SMD (95% CI) |
| Mishel et al, 2009  | PC     | Knowledge              | 20-item question- | 4 weeks post- | 1) Treatment supplemented: Communication strategy DVD + Information Booklet +4 Telephone calls by same nurse to patients and PSP | 15.02 (3.00) | 89 | Handout on staying healthy during treatment | 13.88 (3.83) | 74 | 0.33 (0.02 to 0.64) |
| Mishel et al, 2009  | PC     | Knowledge              | 20-item question- | 3 months post- | 1) Treatment supplemented: Communication strategy DVD + Information Booklet +4 Telephone calls by same nurse to patients and PSP | 15.21 (2.48) | 89 | Handout on staying healthy during treatment | 14.51 (3.32) | 74 | 0.24 (-0.07 to 0.55) |
| Mishel et al, 2009  | PC     | Knowledge              | 20-item question- | 4 weeks post- | 2) Treatment direct: Communication strategy DVD + Information Booklet +4 Telephone calls by nurse | 15.02 (3.00) | 93 | Handout on staying healthy during treatment | 13.88 (3.83) | 74 | 0.33 (0.03 to 0.64) |
| Mishel et al, 2009  | PC     | Knowledge              | 20-item question- | 3 months post- | 2) Treatment direct: Communication strategy DVD + Information + 4 Telephone calls by nurse | 15.51 (2.36) | 93 | Handout on staying healthy during treatment | 14.51 (3.32) | 74 | 0.35 (0.04 to 0.66) |
| Mishel et al, 2009  | PC     | Knowledge              | 20-item question- | 4 weeks post- | 1) +2) | 15.02 (3.00) | 182 | Handout on staying healthy during treatment | 13.88 (3.83) | 74 | 0.35 (0.08 to 0.62) |

**AFFECTIVE COGNITIVE OUTCOMES**

**Knowledge**

| Mishel et al, 2009  | PC     | Knowledge              | 20-item question- | 4 weeks post- | 1) Treatment supplemented: Communication strategy DVD + Information Booklet +4 Telephone calls by same nurse to patients and PSP | 15.02 (3.00) | 89 | Handout on staying healthy during treatment | 13.88 (3.83) | 74 | 0.33 (0.02 to 0.64) |
| Mishel et al, 2009  | PC     | Knowledge              | 20-item question- | 3 months post- | 1) Treatment supplemented: Communication strategy DVD + Information Booklet +4 Telephone calls by same nurse to patients and PSP | 15.21 (2.48) | 89 | Handout on staying healthy during treatment | 14.51 (3.32) | 74 | 0.24 (-0.07 to 0.55) |
| Mishel et al, 2009  | PC     | Knowledge              | 20-item question- | 4 weeks post- | 2) Treatment direct: Communication strategy DVD + Information Booklet +4 Telephone calls by nurse | 15.02 (3.00) | 93 | Handout on staying healthy during treatment | 13.88 (3.83) | 74 | 0.33 (0.03 to 0.64) |
| Mishel et al, 2009  | PC     | Knowledge              | 20-item question- | 3 months post- | 2) Treatment direct: Communication strategy DVD + Information + 4 Telephone calls by nurse | 15.51 (2.36) | 93 | Handout on staying healthy during treatment | 14.51 (3.32) | 74 | 0.35 (0.04 to 0.66) |
| Mishel et al, 2009  | PC     | Knowledge              | 20-item question- | 4 weeks post- | 1) +2) | 15.02 (3.00) | 182 | Handout on staying healthy during treatment | 13.88 (3.83) | 74 | 0.35 (0.08 to 0.62) |

(Continued)
| Author, year | Outcome | Measurement method/tool | Measurement point | Intervention | Control | Effect estimate |
|--------------|---------|-------------------------|-------------------|--------------|---------|-----------------|
| **Mishel et al, 2009**<sup>24</sup> | PC knowledge | 20-item questionnaire (true/false/do not know)<sup>9</sup> | 3 months post-entry after patients had started their treatment | 1) +2) | 15.36 (2.42) | 182 | Handout on staying healthy during treatment | 14.51 (3.32) | 74 | 0.31 (0.04 to 0.58) |
| **Davison et al, 2007**<sup>21</sup> | satisfaction with preparation for treatment decision making with respect to information received | 10-item questionnaire based on 5-point Likert scale<sup>6</sup> | 4–10 weeks after visit - after men had made a treatment decision with/out their Urologist | Generic information 24-min videotape with section on the identification of men’s preferences + Information printout individualized according to patient’s disease characteristics + Written information package with list of relevant info and help sources + Telephone calls by RN approx. 4 weeks after visits | 2.8 (0.84) | 149 | Generic information 24-min videotape + Written information package with list of relevant info and help sources | 2.67 (0.71) | 145 | 0.17 (~0.06 to 0.4) |
| **Hack et al, 2007**<sup>23</sup> | perception of having been informed (total score) | 9-item measure of patient satisfaction with communication with oncologist<sup>5</sup> | 12 weeks post-consultation | Audiotape: audio recording of clinical encounter | 21.83 (3.78) | 214 | Consultation not audiotaped | 21.01 (4.31) | 211 | 0.20 (0.01 to 0.39) |
| **Hack et al, 2007**<sup>23</sup> | perception of having been informed about treatment alternatives | 9-item measure of patient satisfaction with communication with oncologist<sup>5</sup> | 12 weeks post-consultation | Audiotape: audio recording of clinical encounter | 4.73 (0.74) | 214 | Consultation not audiotaped | 4.56 (1.03) | 211 | 0.19 (~0.00 to 0.38) |

*Table 3 (Continued).*
Table 3 (Continued).

| Author, year | Outcome | Measurement method/tool | Measurement point | Intervention | Control | Effect estimate |
|--------------|---------|-------------------------|-------------------|-------------|---------|-----------------|
| Hack et al, 2007 | perception of having been informed about side effects of treatment | 9-item measure of patient satisfaction with communication with oncologist | 12 weeks post-consultation | Audiotape: audio recording of clinical encounter | 4.63 (0.87) 214 | Consultation not audiotaped | 4.37 (1.17) 211 | 0.25 (0.06 to 0.44) |

Mood State/Disturbance

| Mishel et al, 2009 | mood disturbance caused by the personal situation, the disease itself or treatment | 35-item POMS-SF scale | 4 weeks post-entry when treatment decision making meeting had occurred | 1) Treatment supplemented: Communication strategy DVD + Information Booklet +4 Telephone calls by same nurse to patients and PSP | 33.11 (23.29) 89 | Handout on staying healthy during treatment | 37.04 (24.64) 74 | −0.17 (−0.48 to 0.14) |

| Mishel et al, 2009 | mood disturbance caused by the personal situation, the disease itself or treatment | 35-item POMS-SF scale | 3 months post-entry after patients had started their treatment | 1) Treatment supplemented: Communication strategy DVD + Information Booklet +4 Telephone calls by same nurse to patients and PSP | 30.00 (20.4) 89 | Handout on staying healthy during treatment | 32.85 (24.98) 74 | −0.13 (−0.44 to 0.18) |

| Mishel et al, 2009 | mood disturbance caused by the personal situation, the disease itself or treatment | 35-item POMS-SF scale | 4 weeks post-entry when treatment decision making meeting had occurred | 2) Treatment direct: Communication strategy DVD + Information Booklet +4 Telephone calls by nurse | 33.11 (23.29) 93 | Handout on staying healthy during treatment | 37.04 (24.64) 74 | −0.17 (−0.47 to 0.14) |
| Author, year | Outcome | Measurement method/tool | Measurement point | Intervention | Control | Effect estimate |
|-------------|---------|-------------------------|-------------------|--------------|---------|-----------------|
| Mishel et al, 2009<sup>24</sup> | mood disturbance caused by the personal situation, the disease itself or treatment | 35-item POMS-SF scale | 3 months post-entry after patients had started their treatment | 2) Treatment direct: Communication strategy DVD + Information Booklet +4 Telephone calls by nurse | 30.44 (22.06) | 93 Handout on staying healthy during treatment | 32.85 (24.98) | 74 | -0.11 (-0.41 to 0.20) |
| Davison et al, 2007<sup>22</sup> | decisional conflict | 10-item low-literacy scale†† | 4–10 weeks after visit - after men had made a treatment decision with/out their Urologist | Generic information 24-min videotape with section on the identification of men’s preferences + Information printout individualized according to patient’s disease characteristics + Written information package with list of relevant info and help sources + Telephone calls by RN approx. 4 weeks after visits | 1.61 (0.33) | 148 | 1.62 (0.23) | 145 | -0.04 (-0.26 to 0.19) |
| Mishel et al, 2009<sup>24</sup> | decisional regret | 3-item subscale of the QoL scale‡‡ | 3 months post-entry after patients had started their treatment | 1) Treatment supplemented: Communication strategy DVD + Information Booklet +4 Telephone calls by same nurse to patients and PSP | 3.93 (1.82) | 89 | Handout on staying healthy during treatment | 4.17 (1.96) | 74 | -0.13 (-0.44 to 0.18) |
| Mishel et al, 2009<sup>24</sup> | decisional regret | 3-item subscale of the QoL scale‡‡ | 3 months post-entry after patients had started their treatment | 2) Treatment direct: Communication strategy DVD + Information Booklet +4 Telephone calls by nurse | 3.83 (1.44) | 93 | Handout on staying healthy during treatment | 4.17 (1.96) | 74 | -0.2 (-0.51 to 0.11) |
Table 3 (Continued).

| Author, year | Outcome | Measurement method/tool | Measurement point | Intervention | Control | Effect estimate |
|--------------|---------|-------------------------|-------------------|-------------|---------|----------------|
| Mishel et al, 2009 | decisional regret | 3-item subscale of the QoL scale | 3 months post-entry after patients had started their treatment | 1) +2) | 3.88 (1.64) | 182 | Handout on staying healthy during treatment | 4.17 (1.96) | 74 | −0.17 (−0.44 to 0.1) |

**HEALTH OUTCOMES**

**Quality of Life**

Mishel et al, 2009

| Outcome | Measurement method/tool | Measurement point | Intervention | Control | Effect estimate |
|---------|-------------------------|-------------------|-------------|---------|----------------|
| men's perceptions of their overall QoL | 10-item VAS§§ | 4 weeks post-entry when treatment decision making meeting had occurred | 1) Treatment supplemented: Communication strategy DVD + Information Booklet +4 Telephone calls by same nurse to patients and PSP | 7.61 (1.79) | 89 | Handout on staying healthy during treatment | 7.33 (2.23) | 74 | 0.50 (0.18 to 0.81) |

Mishel et al, 2009

| Outcome | Measurement method/tool | Measurement point | Intervention | Control | Effect estimate |
|---------|-------------------------|-------------------|-------------|---------|----------------|
| men's perceptions of their overall QoL | 10-item VAS§§ | 3 months post-entry after patients had started their treatment | 1) Treatment supplemented: Communication strategy DVD + Information Booklet +4 Telephone calls by same nurse to patients and PSP | 7.57 (1.73) | 89 | Handout on staying healthy during treatment | 7.72 (1.72) | 74 | 0.00 (−0.31 to 0.31) |

Mishel et al, 2009

| Outcome | Measurement method/tool | Measurement point | Intervention | Control | Effect estimate |
|---------|-------------------------|-------------------|-------------|---------|----------------|
| men's perceptions of their overall QoL | 10-item VAS§§ | 4 weeks post-entry when treatment decision making meeting had occurred | 2) Treatment direct: Communication strategy DVD + Information Booklet +4 Telephone calls by nurse | 7.61 (1.79) | 93 | Handout on staying healthy during treatment | 7.33 (2.23) | 74 | 0.50 (0.19 to 0.81) |

Mishel et al, 2009

| Outcome | Measurement method/tool | Measurement point | Intervention | Control | Effect estimate |
|---------|-------------------------|-------------------|-------------|---------|----------------|
| men's perceptions of their overall QoL | 10-item VAS§§ | 3 months post-entry after patients had started their treatment | 2) Treatment direct: Communication strategy DVD + Information Booklet +4 Telephone calls by nurse | 7.76 (1.56) | 93 | Handout on staying healthy during treatment | 7.72 (1.72) | 74 | −0.15 (−0.45 to 0.16) |

Notes: 1 Scale: 0 = not at all, 5 = a great deal; items scored and scores converted to 0–100; higher scores = higher levels of satisfaction. 2 Scores: 1 = strongly disagree, 5 = strongly agree; higher scores = higher levels of satisfaction. 3 PC knowledge based on uncertainty management measures. Score based on the sum of the number of correct items. 4 Highest possible satisfaction = score 9. 5 Scores: yes = 1.5, no = 4.5, unsure = 3; range of scores: 15–45; higher scores = higher levels of decision conflict. 6 Higher = greater use of problem solving. 7 Indexed subjects' perceptions of their overall QoL. Scores: 1–10 for how positive men felt about their QoL; higher scores = greater QoL. 8 Abbreviations: SDM, Shared Decision Making; QoL, Quality of Life; POMS-SF, Profile Of Mood States-Short Form; VAS, Visual Analogue Scale; PSP, Primary Supporting Person (eg, wife); RN, Research Nurse; n, number of patients with outcome; N, total number of patients in a group; RR, Relative Risk; SMD, Standard Mean Difference; CI, Confidence Intervals.
estimates showed no significant differences between SDM and usual care on patient satisfaction, decisional conflict, decisional regret, and mood.

This systematic review is based on published data. Surprisingly, six of the thirteen outcomes of interest were not reported. Of particular concern is the dearth of outcome data on the adherence to treatment and health outcomes including symptoms, mortality and QoL. The methodological quality of studies is low to moderate at best, mainly due to unclear allocation concealment, blinding and attrition. The interventions varied in characteristics and content, and were delivered before decision making mostly by trained nurses or research staff. The description of population characteristics was often incomplete too. The studies were carried out mostly in specialized and hospital care in high-income countries from North America, mainly Canada. The ethnicity of the populations was generally not described. Thus, the evidence is generalizable mostly to middle-aged men of at least fifty-five years of age, from Western countries, married, with low to moderate levels of education, who have an English-speaking background and face decision making for PC treatment.

SDM is highly recommended by major task force associations, policy makers, and clinical guidelines for medical decisions regarding PC treatment.\(^{10,11,26–30}\) Our systematic review, however, reveals a lack and variable reporting of outcome measures in the included studies, particularly on health outcomes. This makes it difficult to relate patient involvement in decision making to the actual effects of SDM. The inconsistent reporting of outcomes across studies is most likely due to the lack of consensus in the definition and in the approach used to evaluate and implement SDM.\(^{14}\) SDM is variably adopted in practice as suggested by the very few studies meeting the key SDM criteria. Perspectives of the health care team involved in SDM may play a role in this variability. In a multidisciplinary survey study, 63–71% of oncology nurses, urologists, and oncologists agreed that patients should be involved in decision making.\(^{31}\) However, 52–55% of the urologists and oncologists felt inadequately trained to apply SDM in clinical practice. On the other hand, 20% of the oncology nurses felt inadequately trained to apply SDM. Although an assessment of economic outcomes was outside the focus of our review, we noted that no study reported on cost data.

**Clinical implications**

The variation in the definitions and goals of SDM\(^{14}\) and the inconsistent reporting of outcomes across studies are

**Future research**

Future research warrants further focus on the use of SDM interventions for decisions about PC treatment. Some guidance for SDM implementation based on a clear definition and objectives of SDM as previously reported\(^{14}\) could be the leading step in building focused and solid evidence on SDM. Agreement on a standard set of outcomes that are best to assess SDM and that are most meaningful to patients is critical to guide appropriate outcome collection and evaluation of SDM. Future research then needs to address the absence of evidence, particularly on health outcomes. The link between patient involvement in decision making and the effect of SDM-chosen treatment on patient-important and health-related outcomes needs special consideration. Since treatment can have a significant impact on the patients’ QoL and length of survival, studies of SDM for PC need to provide an appropriate description of the characteristics of the populations including comorbidities, status of cognitive function, literacy levels, sexual health, religious beliefs, side effects of treatment, and whether significant persons (eg, carer, partners) accompany the patients in the process of care. These factors influence not only the type of treatment of choice but also the patients’ emotions, decisional regret, and the degree of involvement in decision making.\(^{32}\) Patients with PC aged <40 years, for example, express significantly high positive and negative emotions, and partners of PC patients express more negative emotions than the patient himself. The physicians’ specialty may also unintentionally influence the physicians’ preference for treatment. Future studies should consider and address the perspectives of the health care team on SDM as a potential barrier for its implementation.\(^{31}\) In particular, interventions and components should be described in detail so that results are interpreted appropriately, and
Interventions can be replicated and evaluated. SDM should also be evaluated in relation to its costs and health gains so that SDM interventions can be reliably implemented. Therefore, future research needs to address the absence of evidence on cost data.

**Strengths and limitations**

To our knowledge, this is the first systematic review on SDM compared to usual care for men faced with decisions about PC treatment. It benefits from the inclusion of international literature without restrictions on countries or type of health care professionals. Including RCTs allows the estimation of causal effects with lower risk of bias. We only included studies meeting the key features of SDM because of the continuing gaps in the conceptualization and implementation of SDM. We considered studies regardless of whether a specific decision was promoted. This was a rigorous and focused approach although we cannot exclude the possibility of underreported SDM characteristics in other studies. We only included literature published in English, but we made considerable efforts to identify all relevant studies. We searched several sources and contacted, between 2015 and 2017, the authors of relevant abstracts with no available full-text, thus increasing the likelihood of identifying more recent literature. The results from our systematic review are limited by the quantity and methodological quality of the available literature, in particular by a dearth of outcome data, and by lower quality reporting of outcome data. The studies focus primarily on SDM process-related outcomes and lack important data on health outcomes, affecting the ability to conduct meta-analyses.

**Conclusion**

There is little research currently available to appropriately evaluate the presumed benefits of SDM for decisions about PC treatment when compared to usual care. SDM may improve knowledge and perception of being informed and may have a positive but short-term effect on patient-perceived QoL. The (long-term) effects of SDM are unclear. Future rigorous research needs a consistent and relevant set of outcomes and interventions to assess the effects of SDM. In particular, it should address the absence of evidence and appropriately describe and test a reproducible form of SDM.

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**Ethics**

No ethical approval was required as this study is a systematic review.

**Disclosure**

The authors declare no conflicts of interest in this work.

**References**

1. Ferlay J, Soerjomataram I, Dikshit R, et al. Cancer incidence and mortality worldwide: sources, methods and major patterns in GLOBOCAN 2012. Int J Cancer. 2015;136(5):E359–E386. doi:10.1002/ijc.29210
2. Center MM, Jemal A, Lortet-Tieulent J, et al. International variation in prostate cancer incidence and mortality rates. Eur Urol. 2012;61(6):1079–1092. doi:10.1016/j.eururo.2012.02.054
3. Ilic D, Neuberger MM, Djulbegovic M, Dahm P. Screening for prostate cancer. Cochrane Database Syst Rev. 2013;(1):Cd004720.
4. Donovan JL, Hamdy FC, Lane JA, et al. Patient-Reported Outcomes after monitoring, surgery, or radiotherapy for prostate cancer. N Engl J Med. 2016;375(15):1425–1437. doi:10.1056/NEJMoa1606221
5. Hamdy FC, Donovan JL, Lane JA, et al. 10-year outcomes after monitoring, surgery, or radiotherapy for localized prostate cancer. N Engl J Med. 2016;375(15):1414–1424. doi:10.1056/NEJMoa1606220
6. Chen R, Clark JA, Talcott JA. Individualizing quality-of-life outcomes reporting: how localized prostate cancer treatments affect patients with different levels of baseline urinary, bowel, and sexual function. J Clin Oncol. 2009;27(24):3916–3922. doi:10.1200/JCO.2008.18.6486
7. Stiggelbout AM, Van der Weijden T, De Wit MP, et al. Shared decision making: really putting patients at the centre of healthcare. BMJ. 2012;344:e256. doi:10.1136/bmj.e256
8. Elwyn G, Frosch D, Thomson R, et al. Shared decision making: a model for clinical practice. J Gen Intern Med. 2012;27(10):1361–1367. doi:10.1007/s11606-012-2077-6
9. Charles C, Gafni A, Whelan T. Decision-making in the physician-patient encounter: revisiting the shared treatment decision-making model. Soc Sci Med. 1999;49(5):651–661.
10. Mottet N, Bellmunt J, Bolla M, et al. EAU-ESTRO-SIOG guidelines on prostate cancer. Part 1: screening, diagnosis, and local treatment with curative intent. Eur Urol. 2017;71(4):618–629. doi:10.1016/j.eururo.2016.08.003
11. Parker C, Gillessen S, Heidenreich A, Horwich A. Cancer of the prostate: ESMO Clinical Practice Guidelines for diagnosis, treatment and follow-up. Ann Oncol. 2015;26 Suppl 5:v69–v77. doi:10.1093/annonc/mdv222
12. Graham J, Kirkbride P, Cann K, Hasler E, Prettyjohns M. Prostate cancer: summary of updated NICE guidance. BMJ. 2014;348:f7524. doi:10.1136/bmj.f7524
13. Xiong T, Turner RM, Wei Y, Neal DE, Lyrratzopoulos G, Higgins J. Comparative efficacy and safety of treatments for localised prostate cancer: an application of network meta-analysis. BMJ Open. 2014;4(5):e004285. doi:10.1136/bmjopen-2013-004285
14. Martinez-González NA, Plate A, Senn O, Markun S, Rosemann T, Neuner-Jehle S. Shared decision-making for prostate cancer screening and treatment: a systematic review of randomised controlled trials. Swiss Med Wkly. 2018;148:w14584. doi:10.4414/smw.2018.14575

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15. Egger M, Smith GD, Altman DG. Principles of and procedures for systematic reviews. In: Egger M, Smith GD, Altman DG, editors. Systematic Reviews in Health Care. 2nd ed. London (UK): BMJ Publishing Group; 2008:23–42.

16. Higgins JPT Green S, editors. Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0. The Cochrane Collaboration; 2009 [updated September 2011]. Available from: www.cochrane-handbook.org. Accessed March, 2018.

17. Liberati A, Altman DG, Tetzlaff J, et al. The PRISMA statement for reporting systematic reviews and meta-analyses of studies that evaluate health care interventions: explanation and elaboration. PLoS Med. 2009;6(7):e1000100. doi:10.1371/journal.pmed.1000100

18. Charles C, Gafni A, Whelan T. Shared decision-making in the medical encounter: what does it mean? (or it takes at least two to tango). Soc Sci Med. 1997;44(5):681–692.

19. Shay LA, Lafata JE. Where is the evidence? A systematic review of shared decision making and patient outcomes. Med Decis Making. 2015;35(1):114–131. doi:10.1177/0272989X14551638

20. Higgins JP, Altman DG, Gotzsche PC, et al. The Cochrane Collaboration’s tool for assessing risk of bias in randomised trials. BMJ. 2011;343:d5928. doi:10.1136/bmj.d5928

21. Review Manager (RevMan) [Computer Program]. Version 5.3.5. Copenhagen: The Nordic Cochrane Centre, The Cochrane Collaboration; 2014. Available from: http://community.cochrane.org/

22. Davison BJ, Goldenberg SL, Wiens KP, Gleave ME. Comparing a generic and individualized information decision support intervention for men newly diagnosed with localized prostate cancer. Cancer Nurs. 2007;30(5):E7–E15. doi:10.1097/01.NCC.0000290819.22195.d6

23. Hack TF, Pickles T, Bultz BD, Ruether JD, Degner LF. Impact of providing audiotapes of primary treatment consultations to men with prostate cancer: a multi-site, randomized, controlled trial. Psychooncology. 2007;16(6):543–552. doi:10.1002/pon.1094

24. Mishel MH, Germino BB, Lin L, et al. Managing uncertainty about treatment decision making in early stage prostate cancer: a randomized clinical trial. Patient Educ Couns. 2009;77(3):349–359. doi:10.1016/j.pec.2009.09.009

25. Davison BJ, Degner LF. Empowerment of men newly diagnosed with prostate cancer. Cancer Nurs. 1997;20(3):187–196.

26. Grossman DC, Curry SJ, Owens DK, et al. Screening for prostate cancer: US preventive services task force recommendation statement. JAMA. 2018;319(18):1901–1913. doi:10.1001/jama.2018.3710

27. Carlsson S, Leapman M, Carroll P, et al. Who and when should we screen for prostate cancer? Interviews with key opinion leaders. BMC Med. 2015;13:288. doi:10.1186/s12916-015-0526-x

28. Bell N, Connor Gorber S, Shane A, et al. Recommendations on screening for prostate cancer with the prostate-specific antigen test. Can Med Assoc J. 2014;186(16):1225–1234. doi:10.1503/cmaj.140703

29. Coulter A, Parsons S, Askham J. Health Systems and Policy Analysis. Policy Brief: Where are the Patients in Decision-Making about Their Own Care? Copenhagen (Denmark): European Observatory on Health Systems and Policies, WHO Regional Office for Europe; 2008.

30. Carter HB, Albertsen PC, Barry MJ, et al. Early detection of prostate cancer: AUA guideline. J Urol. 2013;190(2):419–426. doi:10.1016/j.juro.2013.04.119

31. de Angst IB, Kil PJM, Bangma CH, Takkenberg JJM. Should we involve patients more actively? Perspectives of the multidisciplinary team on shared decision-making for older patients with metastatic castration-resistant prostate cancer. J Geriatr Oncol. 2019. doi:10.1016/j.jgo.2018.12.003

32. Nazim SM, Fawzy M, Bach C, Ather MH. Multi-disciplinary and shared decision-making approach in the management of organ-confined prostate cancer. Arab J Urol. 2018;16(4):367–377. doi:10.1016/j.aju.2018.06.008

33. Elwyn G, Frosch D, Rollnick S. Dual equipoise shared decision making: definitions for decision and behaviour support interventions. Implement Sci. 2009;4(1):75. doi:10.1186/1748-5908-4-75
## Supplementary materials

### Table S1 PRISMA checklist for the reporting of the systematic review

| Section/topic   | # | Checklist item                                                                 | Reported on page # |
|-----------------|---|---------------------------------------------------------------------------------|--------------------|
| **TITLE**       |   |                                                                                  |                    |
| Title           | 1 | Identify the report as a systematic review, meta-analysis, or both.              | 1                  |
| **ABSTRACT**    |   |                                                                                  |                    |
| Structured summary | 2 | Provide a structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number. | 1                  |
| **INTRODUCTION**|   |                                                                                  |                    |
| Rationale       | 3 | Describe the rationale for the review in the context of what is already known.   | 1-2                |
| Objectives      | 4 | Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS). | 2                  |
| **METHODS**     |   |                                                                                  |                    |
| Protocol and registration | 5 | Indicate if a review protocol exists, if and where it can be accessed (eg, Web address), and, if available, provide registration information including registration number. | 2                  |
| Eligibility criteria | 6 | Specify study characteristics (eg, PICOS, length of follow-up) and report characteristics (eg, years considered, language, publication status) used as criteria for eligibility, giving rationale. | 2                  |
| Information sources | 7 | Describe all information sources (eg, databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched. | 2                  |
| Search          | 8 | Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated. | Table S2           |
| Study selection | 9 | State the process for selecting studies (ie, screening, eligibility, included in systematic review, and, if applicable, included in the meta-analysis). | 2                  |
| Data collection process | 10 | Describe method of data extraction from reports (eg, piloted forms, independently, in duplicate) and any processes for obtaining and confirming data from investigators. | 2                  |
| Data items      | 11 | List and define all variables for which data were sought (eg, PICOS, funding sources) and any assumptions and simplifications made. | 2                  |
| Risk of bias in individual studies | 12 | Describe methods used for assessing risk of bias of individual studies (including specification of whether this was done at the study or outcome level), and how this information is to be used in any data synthesis. | 2                  |
| Summary measures | 13 | State the principal summary measures (eg, risk ratio, difference in means). | 2                  |
| Synthesis of results | 14 | Describe the methods of handling data and combining results of studies, if done, including measures of consistency (eg, I²) for each meta-analysis. | 2                  |
| Risk of bias across studies | 15 | Specify any assessment of risk of bias that may affect the cumulative evidence (eg, publication bias, selective reporting within studies). | 2                  |
| Additional analyses | 16 | Describe methods of additional analyses (eg, sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified. | 2                  |

(Continued)
Table S1 (Continued).

| Section/topic          | #  | Checklist item                                                                 | Reported on page # |
|------------------------|----|---------------------------------------------------------------------------------|--------------------|
| **RESULTS**            |    |                                                                                 |                    |
| Study selection        | 17 | Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a flow diagram. | 3, Figure 1        |
| Characteristics        | 18 | For each study, present characteristics for which data were extracted (eg, study size, PICOS, follow-up period) and provide the citations. | 3-4, Table 1       |
| Risk of bias within studies | 19 | Present data on risk of bias of each study and, if available, any outcome level assessment (see item 12). | 4, Table 2         |
| Results of individual studies | 20 | For all outcomes considered (benefits or harms), present, for each study: (a) simple summary data for each intervention group (b) effect estimates and confidence intervals, ideally with a forest plot. | 4, 9, Figure 2, Table 3 |
| Synthesis of results   | 21 | Present results of each meta-analysis done, including confidence intervals and measures of consistency. | NA                 |
| Risk of bias across studies | 22 | Present results of any assessment of risk of bias across studies (see Item 15). | Table 2            |
| Additional analysis    | 23 | Give results of additional analyses, if done (eg, sensitivity or subgroup analyses, meta-regression [see Item 16]). | NA                 |
| **DISCUSSION**         |    |                                                                                 |                    |
| Summary of evidence    | 24 | Summarize the main findings including the strength of evidence for each main outcome; consider their relevance to key groups (eg, health care providers, users, and policy makers). | 9, 16              |
| Limitations            | 25 | Discuss limitations at study and outcome level (eg, risk of bias), and at review-level (eg, incomplete retrieval of identified research, reporting bias). | 17                 |
| Conclusions            | 26 | Provide a general interpretation of the results in the context of other evidence, and implications for future research. | 17                 |
| **FUNDING**            |    |                                                                                 |                    |
| Funding                | 27 | Describe sources of funding for the systematic review and other support (eg, supply of data); role of funders for the systematic review. | No external funding |

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Table S2 Search strategy for OVID Medline

| Item | Searches |
|------|----------|
| 1    | exp Decision Making/or Decision Making, Organizational/or Decision Trees/or Decision Making/or Decision Support Techniques/or Decision Support Systems, Clinical/or Decision Making, Computer-Assisted/or exp Computer-Assisted Instruction/or exp Patient Participation/or exp Professional-Patient Relations/or exp “Attitude of Health Personnel”/or Counseling/or exp Health Communication/ |
| 2    | exp Informed Consent/ |
| 3    | (choice behavior or decision making or shared decision making).mp.tw. |
| 4    | (informed adj3 (consent or choice or decision*)).mp.tw. |
| 5    | ((decision* or decid*) adj4 (support* or aid* or tool* or instrument* or technolog* or technique* or system* or program* or algorithm* or process* or method* or intervention* or material*)).mp.tw. |

(Continued)
Table S2 (Continued).

| Item | Searches |
|------|----------|
| 6    | (decision adj3 (board* or guide* or counseling)).mp,tw. |
| 7    | (computer* adj4 decision making).mp. |
| 8    | (patient adj3 (participation or involvement or cent#d care)).mp,tw. |
| 9    | ((risk communication or risk assessment or risk information) adj4 (tool* or method*)).mp,tw. |
| 10   | interact* health communication*.mp,tw. |
| 11   | (interact* adj (internet or online or graphic* or booklet*)).mp,tw. |
| 12   | (interact* adj4 tool*).mp,tw. |
| 13   | ((interact* or evidence based) adj3 (risk information or risk communication or risk presentation or risk graphic*)).mp,tw. |
| 14   | adaptive conjoint analys#s.mp,tw. |
| 15   | or/1–14 |
| 16   | (Prostat* adj3 (Neoplasm* or Cancer or tumo?r* or carcinoma)).mp,tw. |
| 17   | exp Prostatic Neoplasms/ |
| 18   | 16 or 17 |
| 19   | 15 and 18 |
| 20   | (letter or letter$ or editorial or historical article or anecdote or commentary or note or case report$ or case study).pt,sh. |
| 21   | (animals not humans).sh. |
| 22   | 20 or 21 |
| 23   | 19 not 22 |
| 24   | exp Randomized Controlled Trial/or exp clinical trial/ |
| 25   | randomized controlled trial.pt. |
| 26   | randomized controlled trial.sh. |
| 27   | controlled clinical trial.pt. |
| 28   | random allocation.sh. |
| 29   | double blind method.sh. |
| 30   | single blind method.sh. |
| 31   | or/24–30 |
| 32   | 31 not 22 |
| 33   | exp clinical trial/or exp Clinical Trials as Topic/ |
| 34   | clinical trial.pt. |
| 35   | ((sing$ or doubl$ or trebl$ or trpl$) adj25 (blind$ or mask$)).ti,ab. |
| 36   | (clin$ adj25 trial$).ti,ab. |
| 37   | (random$ or placebo$).ti,ab. |
| 38   | (PLACEBO or RESEARCH DESIGN).sh. |

(Continued)
**Table S2 (Continued).**

| Item | Searches |
|------|----------|
| 39   | or/33–38 |
| 40   | 39 not 22 |
| 41   | 40 not 32 |
| 42   | exp EVALUATION STUDIES/ |
| 43   | (comparative study or follow up studies or prospective studies).sh. |
| 44   | (control$ or prospectiv$ or volunteer$).ti,ab. |
| 45   | or/42–44 |
| 46   | 45 not 22 |
| 47   | 46 not (32 or 41) |
| 48   | 23 and (32 or 41 or 47) |

**Notes:** *Similar strategies were developed in EMBASE (Elsevier), CINHAL (EBSCOHost), The Cochrane Library (Wiley), PsychINFO (EBSCOHost), and Scopus.

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**Reference**

1. Moher D, Liberati A, Tetzlaff J, Altman DG, The PRISMA Group (2009) Preferred Reporting Items for Systematic Reviews and Meta-Analyses: The PRISMA Statement. PLoS Med 6(7): e1000097. doi:10.1371/journal.pmed.1000097