

# BMJ Open Coffee consumption and risk of prostate cancer: a systematic review and meta-analysis

Xiaonan Chen, Yiqiao Zhao, Zijia Tao, Kefeng Wang 

**To cite:** Chen X, Zhao Y, Tao Z, *et al.* Coffee consumption and risk of prostate cancer: a systematic review and meta-analysis. *BMJ Open* 2021;**11**:e038902. doi:10.1136/bmjopen-2020-038902

► Prepublication history and additional material for this paper is available online. To view these files, please visit the journal online (<http://dx.doi.org/10.1136/bmjopen-2020-038902>).

Received 11 April 2020

Revised 24 September 2020

Accepted 21 October 2020

## ABSTRACT

**Objectives** To conduct a systematic review with meta-analysis of cohort studies to evaluate the association of coffee consumption with the risk of prostate cancer.

**Data sources** PubMed, Web of Science and Embase were searched for eligible studies up to September 2020.

**Study selection** Cohort studies were included.

**Data extraction and synthesis** Two researchers independently reviewed the studies and extracted the data. Data synthesis was performed via systematic review and meta-analysis of eligible cohort studies. Meta-analysis was performed with the “*metan*” and “*glst*” commands in Stata 14.0.

**Main outcomes and measures** Prostate cancer was the main outcome. It was classified as localised prostate cancer which included localised or non-aggressive cancers; advanced prostate cancer which included advanced or aggressive cancers; or fatal prostate cancer which included fatal/lethal cancers or prostate cancer-specific deaths.

**Results** Sixteen prospective cohort studies were finally included, with 57 732 cases of prostate cancer and 1 081 586 total cohort members. Higher coffee consumption was significantly associated with a lower risk of prostate cancer. Compared with the lowest category of coffee consumption, the pooled relative risk (RR) was 0.91 (95% CI 0.84 to 0.98),  $I^2=53.2\%$  for the highest category of coffee consumption. There was a significant linear trend for the association ( $p=0.006$  for linear trend), with a pooled RR of 0.988 (95% CI 0.981 to 0.995) for each increment of one cup of coffee per day. For localised, advanced and fatal prostate cancer, the pooled RRs were 0.93 (95% CI 0.87 to 0.99), 0.88 (95% CI 0.71 to 1.09) and 0.84 (95% CI 0.66 to 1.08), respectively. No evidence of publication bias was indicated in this meta-analysis.

**Conclusions** This study suggests that a higher intake of coffee may be associated with a lower risk of prostate cancer.

## INTRODUCTION

Prostate cancer is the second most frequently diagnosed cancer and the sixth leading cause of cancer death in men. There were 1 276 000 new cancer cases and 359 000 cancer deaths in 2018.<sup>1</sup> It is estimated that nearly three-quarters of prostate cancer cases occur in developed countries.<sup>1</sup> Since the 1970s, the incidence of prostate cancer has also

## Strengths and limitations of this study

- Risk of selection and recall bias may be minimised due to the inclusion of prospective cohort studies.
- Large sample size ensures adequate statistical power to detect even a small effect of interest.
- Uncontrolled/residual confounding may distort the association between coffee consumption and prostate cancer.
- Misclassification of coffee consumption may occur due to the self-reported nature of the exposure.
- Significant heterogeneity among study results may come from various sources.

increased rapidly in some Asian countries such as China, Singapore and Japan, where the incidence has always been much lower than in some Western countries.<sup>1 2</sup> Therefore, primary prevention of prostate cancer is a significant public health problem worldwide.

Coffee is one of the most popular beverages. Since its popularity continues to increase worldwide, even a small effect on individual health may exert a substantial public health impact. Coffee is known to be a major source of dietary caffeine, cafestol and antioxidants in industrialised nations.<sup>3</sup> Its various constituents such as caffeine, caffeic acid and chlorogenic acid can potentially impact the development of cancer through multiple carcinogenesis pathways.<sup>4 5</sup> Inverse associations were observed between coffee consumption and the risk of cancer in sites such as the liver, colorectum and breast.<sup>6</sup> However, previous studies have reported inconsistent results on the association of coffee consumption with the risk of prostate cancer. Although earlier cohort studies did not detect an association,<sup>7–15</sup> more recent studies conducted in major Western countries such as the USA, Sweden and the UK reported that coffee consumption was associated with a lower risk of localised and advanced prostate cancer.<sup>16–20</sup> In Japan, a country with increasing popularity of coffee, a cohort study also found a



© Author(s) (or their employer(s)) 2021. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

Department of Urology, Shengjing Hospital of China Medical University, Shenyang, Liaoning, China

## Correspondence to

Dr Kefeng Wang;  
[wangkefenguro@sina.com](mailto:wangkefenguro@sina.com)

significant inverse association between coffee consumption and the risk of prostate cancer.<sup>21</sup>

Previous meta-analyses of cohort studies up to 2015 reported a significant positive association for coffee consumption on total prostate cancer risk, with highly variable results in different subgroups.<sup>22–23</sup> Since then, five cohort studies have explored the association but still reported inconsistent results.<sup>24–28</sup> It was hypothesised that higher coffee consumption was associated with an increased risk of prostate cancer. Thus, the objective of this updated meta-analysis was to explore and evaluate the association of coffee intake with the risk of prostate cancer in adult men, and to direct the future primary prevention strategy on prostate cancer.

## METHODS

This systematic review was conducted and reported in adherence to the PRISMA and MOOSE guidelines<sup>29</sup>; the corresponding checklists are shown in online supplemental table 1 and online supplemental file 1. Two researchers (YZ and ZT) independently conducted the literature search, study selection, data extraction and study quality assessment. Any discrepancies were resolved by discussion, but whenever consensus could not be reached between the two reviewers, a third reviewer (KW) acted as arbitrator.

### Patient and public involvement

This is a meta-analysis based on study-level data and no individual-level data were involved in the study or in defining the research question or outcome measures.

### Inclusion criteria

The eligibility criteria of the studies were as follows: (1) The study should use a longitudinal cohort design or case-control design nested within a cohort study. (2) The study should present information on coffee consumption as the exposure of interest. Coffee consumption was ascertained by self-reported dietary records or food diaries on the intake levels (highest intake category vs lowest intake category) or frequency measures (eg, per unit/cups/mL per day/week). Since the intake levels were classified and defined differently in each study, the absolute coffee consumption in the highest and lowest intake categories varied across the included studies. (3) The study should report prostate cancer as the outcome of interest. Prostate cancer was defined by clinical diagnosis, physician diagnosis, medical records, self-reports or data linkage to a registry system such as a cancer registry. Based on definitions in each original study, the prostate cancer categories were classified as follows: (a) localised prostate cancer, which included localised or non-aggressive cancers; (b) advanced prostate cancer, which included advanced or aggressive cancers; and (c) fatal prostate cancer, which included fatal/lethal cancers or prostate cancer-specific deaths. (4) The study should provide relative risk (RR), HR, risk ratio, rate ratio or OR estimates with confidence

intervals (CIs) or standard errors for the association of coffee consumption with the risk of prostate cancer. If multiple estimates were provided, priority was given to the multivariable-adjusted risk estimates. If more than one study was conducted in the same population, the earlier reports or reports with less applicable information were excluded.

### Literature search

A literature search was performed using PubMed, Web of Science and Embase up to September 2020 with the following keywords: coffee and prostate and (cancer or carcinoma or neoplasm or tumour). The full search strategy is shown in online supplemental file 2. The reference lists of relevant publications were also manually searched for identification of additional eligible studies. No language limitation was imposed.

When data or information in the publication were insufficient, we attempted to contact the corresponding authors of the original study to request the relevant data. Then two authors (Russnes and Nilsson) provided us with the relevant information about the person-years of follow-up for specific categories of coffee intake to facilitate the dose-response analyses.<sup>15–20</sup> Of note, we finally did not include the study by Russnes *et al*<sup>20</sup> in the current meta-analysis because the study population is the same as another included cohort study by Wilson *et al*<sup>18</sup> which reported more applicable information.

### Data extraction

We extracted the following information from each eligible study: first author's name, year of publication, study country, follow-up time, number of participants in the cohort, number of prostate cancer cases, assessment of coffee consumption, primary study outcome, definitions and categories of coffee consumption, RRs and 95% CIs for all prostate cancer outcomes associated with coffee consumption, and the potential confounders considered or adjusted in the analysis.

### Study quality assessment

The 9-star Newcastle–Ottawa Scale tool was used to assess study quality.<sup>30</sup> The quality of each cohort study was judged on three broad categories—namely, selection of the study population, comparability of groups and ascertainment of either the exposure or outcome of interest.

### Statistical analysis

In the meta-analysis, the RR estimate was used to measure the association between coffee consumption and the risk of prostate cancer in this meta-analysis. We pooled the study-specific RR estimates for the highest versus the lowest category of coffee consumption. A fixed effects model was used to pool the study-specific estimates; whenever significant heterogeneity was detected, the random effect model was used to address the heterogeneity across studies.<sup>31</sup> Subgroup analyses were conducted stratified by study location, prostate cancer stage and potential confounder adjustments including a history of

prostate-specific antigen (PSA) testing, a family history of prostate cancer, total energy intake, cigarette smoking, alcohol consumption, physical activity, body mass index (BMI) or history of diabetes. Since PSA testing was generally introduced after 1986,<sup>32</sup> studies with follow-up periods that ended before 1986 were classified in the PSA-adjusted group. To explore the influence of each study on the pooled results, sensitivity analyses were also performed by excluding one study at a time and then repeating the meta-analysed approach.

We further examined the potential dose-response relationship between coffee consumption and the risk of prostate cancer. When the mean coffee intakes in each category were not reported, the midpoint values in each category were used instead; when the upper boundary of the highest intake category was not presented, we calculated the midpoint value assuming that the highest category had the same magnitude of intake as the preceding category.<sup>33 34</sup> The pooled RR for each increment of one cup of coffee per day was estimated using the method proposed by Orsini and Greenland.<sup>35</sup> We examined a potential non-linear relation between coffee consumption and prostate cancer risk by modelling coffee consumption using restricted cubic splines for non-linear trends with 4 knots at fixed percentiles (5%, 35%, 65% and 95%) of the distribution.<sup>36</sup> Non-linearity of the association was explored by testing the null hypothesis that the coefficients of the second and third splines were equal to zero.

We assessed the heterogeneity by using the Q and the  $I^2$  statistic. A p value <0.10 or an  $I^2$  >50% suggest that statistical heterogeneity may exist.<sup>37</sup> Small study effects such as publication bias were evaluated by funnel plots, as well as Begg's test and Egger's test.<sup>38 39</sup> Meta-analysis was conducted using the "metan" and "glst" commands in Stata version 14.0 (StataCorp, College Station, Texas, USA). Two-sided p values <0.05 were considered statistically significant in the meta-analysis.

## RESULTS

### Literature search

We identified 497 records after searching the three databases. After 217 duplicate records were removed, 280 records remained for screening of titles and abstracts, and after screening the titles and abstracts, 254 irrelevant records were excluded. Following a further full-text review of the 26 remaining studies, 10 studies were excluded because of no useful risk estimates or 95% CIs; two studies were excluded as newer data or more informative data were available. Fourteen studies were obtained from full-text screening and a further two studies were identified by checking the reference lists of retrieved articles.<sup>9 11</sup> Thus, 16 studies were included in the final analysis,<sup>9-11 13-19 21 24-28</sup> of which 15 reported on the risk of prostate cancer associated with the highest versus the lowest coffee consumption<sup>9-11 13-19 21 24-27</sup>; 13 studies reported the risk associated with an increase of one cup

of coffee per day or provided sufficient data to estimate the dose-response risk<sup>9 13-19 21 24 25 27 28</sup> (figure 1).

### Study characteristics and quality assessment

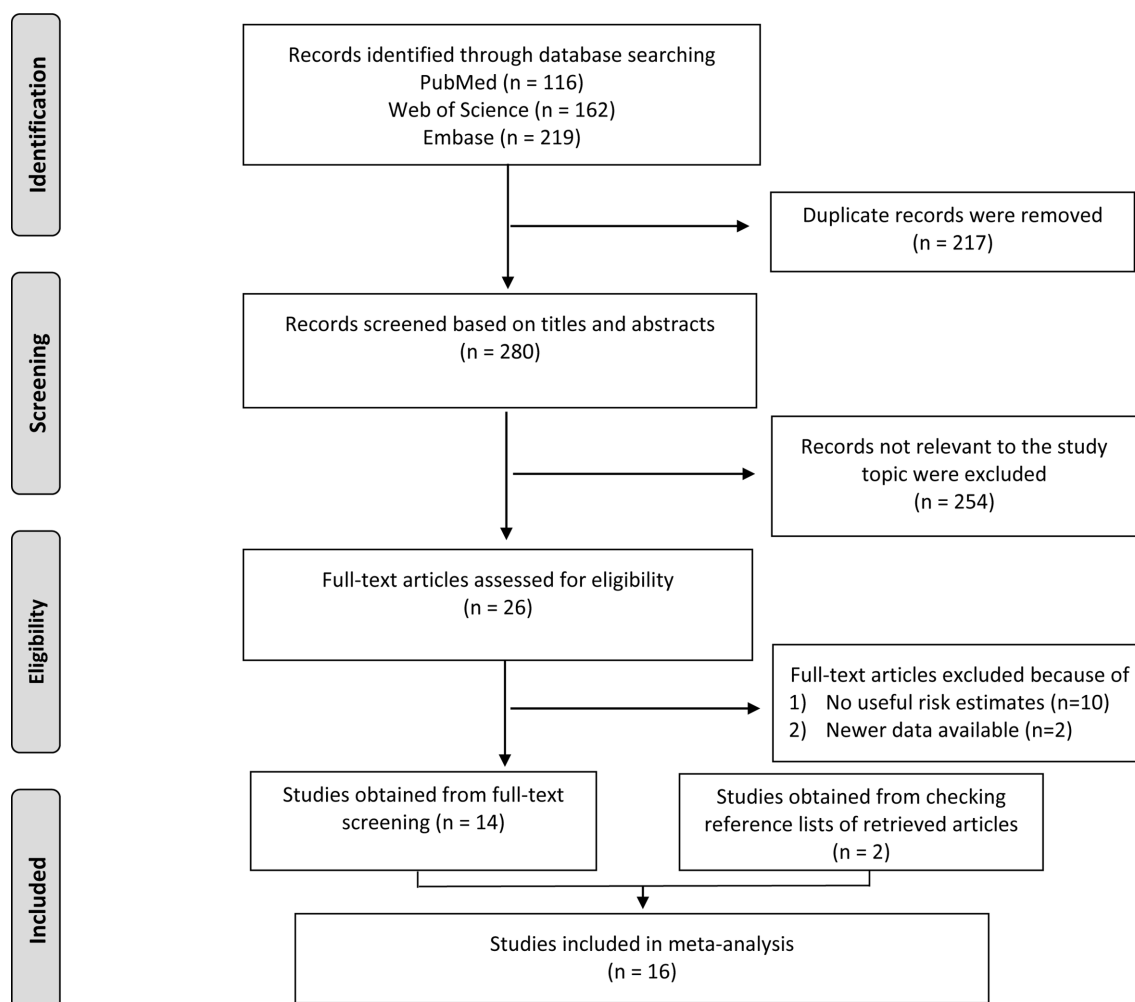
The characteristics of the eligible cohort studies are shown in online supplemental file 3. The included studies were conducted in North America (n=7), Europe (n=7) and Japan (n=2). There was a total of 1 081 586 men in the 16 cohort studies, of whom 57 732 developed prostate cancer. To measure coffee consumption, 11 studies used food-frequency questionnaires and five used a self-administered dietary questionnaire. Most studies considered or adjusted for the most potential confounders in the analysis, such as age at baseline, family history of prostate cancer, race, cigarette smoking, alcohol drinking, total energy intake, BMI and physical activity. The results of study quality assessment are presented in online supplemental file 4. The total scores for each cohort study ranged from 6 to 9. Fourteen studies awarded a total score of ≥7, which were considered as relatively high-quality studies with a low risk of bias.

### Overall analyses and dose-response analyses

The reported RRs for the original cohort studies ranged from 0.47 (95% CI 0.25 to 0.87) in the study by Pounis *et al* to 1.42 (95% CI 0.77 to 2.61) in the study by Ellison *et al* (figure 2). Compared with the lowest coffee intake category, there was a 9% reduction in the risk of prostate cancer for the highest category (RR=0.91; 95% CI 0.84 to 0.98). Statistically significant heterogeneity was detected across the studies (p=0.008,  $I^2$ =53.2%). In dose-response analyses, we found evidence of a linear inverse association between coffee consumption and prostate cancer risk (p=0.006 for linear trend) (figure 3). The pooled RR of prostate cancer was 0.988 (95% CI 0.981 to 0.995) for an increase of one cup of coffee per day. No evidence of a non-linear relationship was observed between coffee consumption and risk of prostate cancer (p=0.193 for non-linearity). Moreover, there was no indication of small study effects such as publication bias from the results of the Egger's test (p=0.409), Begg's test (p=0.843) as well as the funnel plot. The funnel plot and Egger's publication bias plot are shown in online supplemental file 5.

### Subgroup and sensitivity analyses

As shown in table 1, compared with the lowest coffee intake category there was a 7% reduction in risk for the highest intake category (RR=0.93; 95% CI 0.87 to 0.99) for localised prostate cancer. For advanced and fatal prostate cancer, the corresponding pooled RRs were 0.88 (95% CI 0.71 to 1.09) and 0.84 (95% CI 0.66 to 1.08) (figure 4). When stratified by study location, the pooled RRs were 0.96 (95% CI 0.90 to 1.03), 0.85 (95% CI 0.74 to 0.98) and 0.85 (95% CI 0.48 to 1.51) for studies conducted in North America (six in the USA and one in Canada), European countries and Japan. Furthermore, significant inverse associations were observed in all of the confounder adjusted subgroups.



**Figure 1** Flow diagram of study selection in the meta-analysis.

In sensitivity analyses, we sequentially excluded one study at a time and recalculated the pooled RRs of the remaining studies. The pooled RRs did not change substantially, ranging from 0.89 (95% CI 0.82 to 0.97) to 0.93 (95% CI 0.86 to 1.00) after omission of the studies by Hashibe *et al* and Terdal *et al*, respectively.

## DISCUSSION

### Summary of the findings

In this meta-analysis, higher coffee consumption was significantly associated with a reduced risk of prostate cancer in men. In the dose-response analysis, a reduction in the risk of prostate cancer of nearly 1% was observed for each increment of one cup of coffee per day. The combined estimate for prostate cancer was robust across subgroup and sensitivity analyses.

### Comparison with other studies

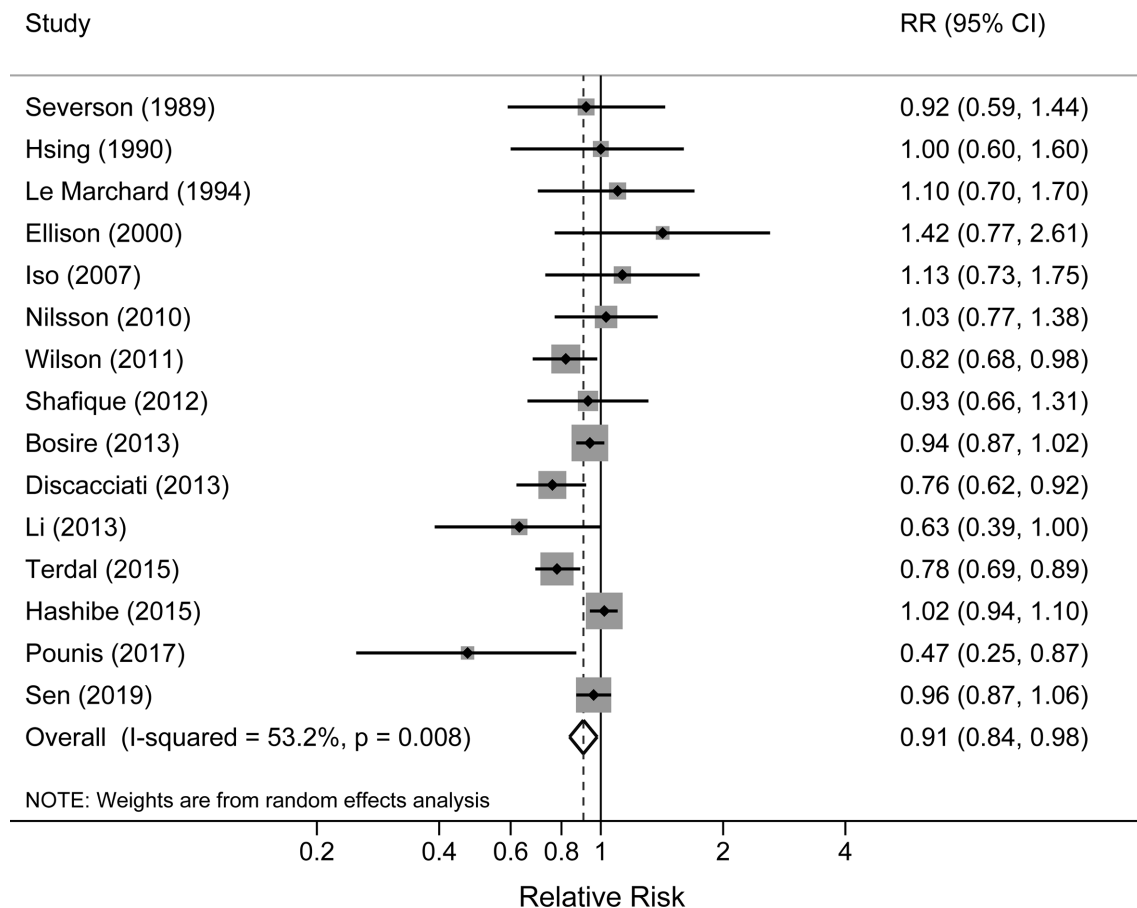
The previous meta-analysis detected a statistically significant positive association between coffee consumption and prostate cancer risk (RR=1.16; 95% CI 1.01 to 1.33).<sup>22</sup> However, this observed effect was confined to case-control studies (RR=1.21; 95% CI 1.03 to 1.43), with no significant

association in cohort studies (RR=1.06; 95% CI 0.83 to 1.35) when stratified by study design.<sup>22</sup> Considering the case-control design, patients with prostate cancer might differentially recall their past coffee consumption habits compared with healthy controls, which might generally lead to biased estimates. This potential recall bias could generate a spurious positive association between coffee consumption and prostate cancer risk. Additionally, selection bias, which can occur in case-control studies, may distort the association between coffee consumption and prostate cancer risk. In another meta-analysis of cohort studies with 539 577 participants and 34 105 prostate cancer cases, the pooled RR for the highest category of coffee intake was 0.90 (95% CI 0.85 to 0.95) for total prostate cancer compared with the lowest intake category. In this updated meta-analysis of 1 081 586 cohort members and 57 732 incident cases, the overall result was similar to the previous one. However, for subgroups of localised, advanced and fatal prostate cancers, the strength of associations tended to be weaker compared with the previous study.

### Possible biological mechanisms

It is biologically plausible that coffee may reduce the risk of prostate cancer in men. Coffee improves

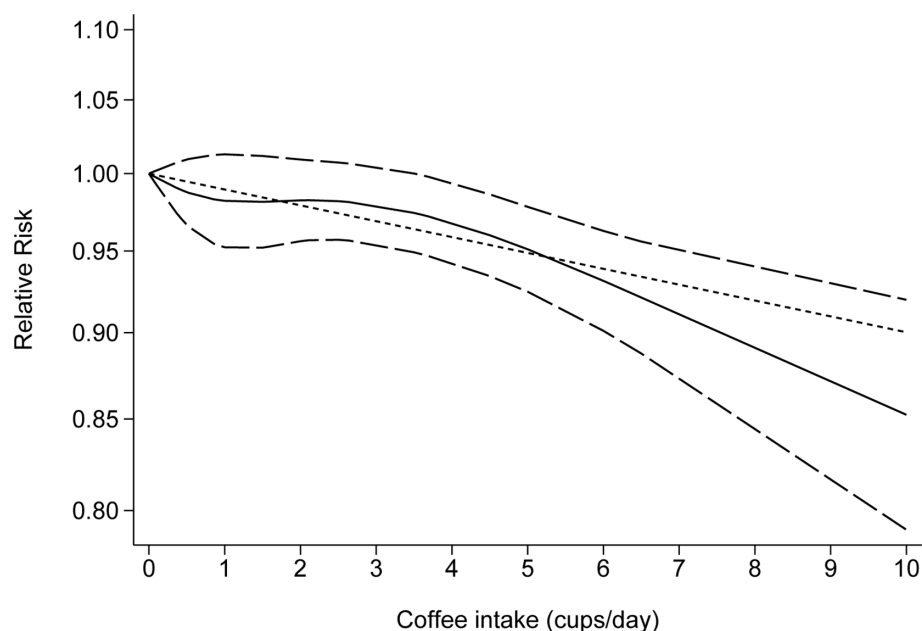




**Figure 2** Forest plot for the association between coffee consumption and prostate cancer risk.

glucose metabolism, decreases concentrations of plasma insulin and insulin-like growth factor-1, has anti-inflammatory and antioxidant effects, and affects sex hormone levels, all of which may play roles in the initiation, development and progression of prostate

cancer.<sup>3 18 20 40</sup> Coffee is also a major source of chlorogenic acids; intake of quinides, the degradation products of chlorogenic acids, has been observed to increase insulin sensitivity and lower blood glucose levels.<sup>3</sup> Moreover, coffee intake may be associated



**Figure 3** Dose-response relationship of coffee consumption with prostate cancer risk.

**Table 1** Summary risk estimates and corresponding 95% CIs for prostate cancer associated with the highest versus the lowest coffee consumption

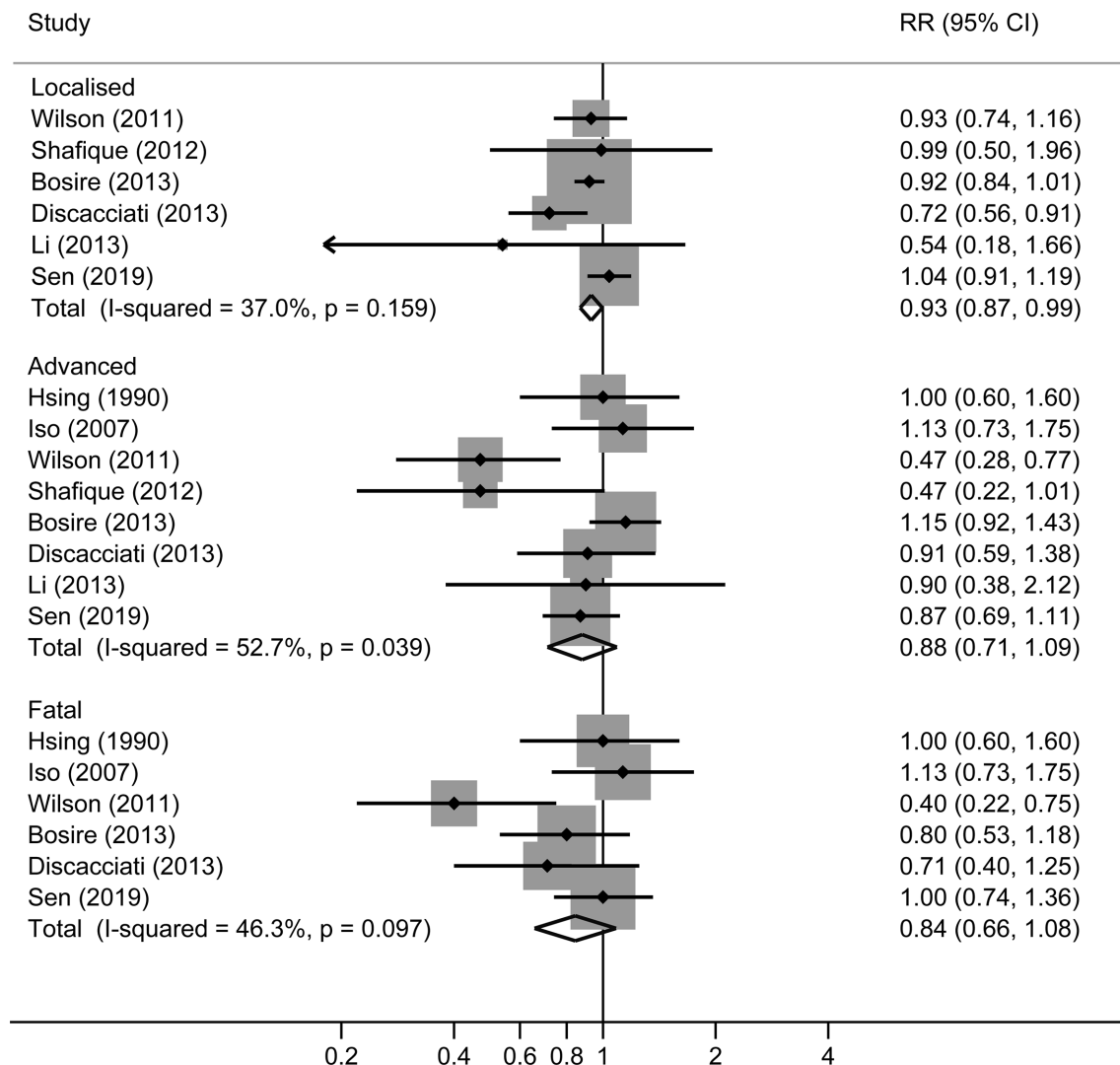
	No of studies	Summary RR	95% CI	I <sup>2</sup> (%)	P value*
Overall	15	0.91	(0.84 to 0.98)	53.2	0.008
Prostate cancer category †					
Localised	6	0.93	(0.87 to 0.99)	37.0	0.159
Advanced	8	0.88	(0.71 to 1.09)	52.7	0.039
Fatal	6	0.84	(0.66 to 1.08)	46.3	0.097
Study location					
North America	7	0.96	(0.90 to 1.03)	17.7	0.295
Europe	6	0.85	(0.74 to 0.98)	63.3	0.018
Japan	2	0.85	(0.48 to 1.51)	68.5	0.075
NOS score					
6	2	1.20	(0.84 to 1.72)	0	0.507
7	3	1.02	(0.78 to 1.32)	0	0.810
8 or 9	10	0.88	(0.81 to 0.96)	66.2	0.002
<b>Adjustment for confounders</b>					
PSA testing ‡					
Yes	6	0.86	(0.77 to 0.96)	31.8	0.197
No	9	0.94	(0.84 to 1.06)	60.5	0.009
Family history of prostate cancer					
Yes	4	0.83	(0.72 to 0.96)	57.8	0.068
No	11	0.95	(0.85 to 1.05)	50.8	0.026
Total energy intake					
Yes	6	0.85	(0.76 to 0.96)	61.1	0.025
No	9	0.97	(0.85 to 1.09)	47.7	0.053
Smoking status					
Yes	10	0.86	(0.79 to 0.94)	52.0	0.027
No	5	1.03	(0.95 to 1.11)	0	0.805
Alcohol consumption					
Yes	6	0.87	(0.84 to 0.98)	49.2	0.008
No	9	0.93	(0.84 to 1.03)	57.2	0.017
Physical activity					
Yes	7	0.87	(0.79 to 0.95)	58.4	0.025
No	8	1.00	(0.90 to 1.12)	10.5	0.348
BMI					
Yes	9	0.86	(0.78 to 0.94)	56.9	0.017
No	6	1.03	(0.95 to 1.11)	0	0.897
Diabetes					
Yes	5	0.87	(0.84 to 0.98)	64.9	0.022
No	10	0.97	(0.86 to 1.10)	22.8	0.233

\*P value for heterogeneity within each subgroup.

†Based on definition in each original study, the prostate cancer categories were classified as follows: (a) localised prostate cancer which included localised or non-aggressive cancers, (b) advanced prostate cancer which included advanced or aggressive cancers, (c) fatal prostate cancer which included fatal/lethal cancers or prostate cancer-specific deaths.

‡Since PSA testing was generally introduced after 1986, studies with follow-up periods that ended before 1986 were classified in the PSA-adjusted group.

BMI, body mass index; NOS, Newcastle–Ottawa Scale; PSA, prostate-specific antigen; RR, relative risk.



**Figure 4** Forest plot for the association between coffee consumption and risk of prostate cancer stratified by cancer stages.

with increased levels of adiponectin plasma,<sup>41 42</sup> which may act as an endogenous insulin sensitiser.<sup>43</sup> Higher adiponectin levels in plasma are supposed to relate to lower concentrations of plasma insulin.<sup>43</sup> In two prospective studies, insulin levels were observed to be directly associated with prostate cancer-specific mortality.<sup>44 45</sup>

Coffee is a major contributor of dietary antioxidants such as caffeic acid and chlorogenic acid.<sup>20</sup> A prospective cohort study from the USA found that dietary antioxidants from coffee (eg, caffeic acid) were inversely associated with the risk of total, advanced and lethal prostate cancer.<sup>20</sup> It was suggested that antioxidants protect cells from damage caused by oxidative stress and inflammation, which may further lead to neoplastic transformation in the prostate.<sup>46</sup> Additionally, dietary antioxidants may inhibit prostate cancer progression through suppression of oxidative stress, which might play a critical role during the progression of prostate cancer.<sup>46</sup> Coffee intake was indicated to be related to increased levels in sex hormone-binding globulin (SHBG), as well as total testosterone.<sup>47 48</sup> A

pooled analysis of 18 prospective studies found that SHBG levels may be inversely associated with the risk of prostate cancer.<sup>40</sup> Of note, a nested case-control study found that caffeine or caffeinated coffee intakes were suggested to be associated with an increased level of plasma SHBG. However, such an association was not observed between decaffeinated coffee and plasma SHBG levels. Thus, it was suggested that caffeine may be the key component in coffee, which may be responsible for determining plasma SHBG levels.<sup>48</sup>

### Strengths and limitations

A strength of this study was the inclusion of prospective cohort studies. Cohort studies could minimise the risk of selection and recall bias, which is a major concern for case-control design. Besides, large numbers of total cohort members and prostate cancer cases ensure adequate statistical power to detect even a small effect of interest. Furthermore, the dose-response analysis may further lend confidence to the study hypothesis that increased coffee consumption was linearly associated with a lower risk of prostate cancer. Besides,

most of the studies were of high quality with a low risk of bias, which could further lend confidence to the current pooled results.

This meta-analysis also has several limitations. First, one weakness is that only three databases were searched for eligible studies and other databases, especially non-English databases, were not considered in the literature search. Second, because of the observational design, unmeasured or uncontrolled confounders in the original studies may bias the pooled risk estimate; however, the residual confounding effects from the original studies were difficult to handle in a meta-analysis approach.<sup>49 50</sup> For example, the inverse association between coffee consumption and prostate cancer could be attributed to risk factors related to coffee consumption, such as physical activity and healthy diet. However, most of the original studies have considered or adjusted for these major potential confounders in the analysis. In the sensitivity analysis of restricting the meta-analysis in studies considering most confounders, the strength of association tended to be larger in comparison with the overall association. Third, misclassification of coffee consumption may occur because of the self-reported nature of exposure measurement. However, validation studies by diet records indicated a relatively high validity of coffee consumption measured by food frequency questionnaire. The correlations between questionnaire and diet records were 0.80 in US men,<sup>16</sup> 0.71 in Swedish men<sup>17</sup> and 0.72 in Japanese men.<sup>21</sup> Of note, misclassification of exposure would most likely be non-differential in cohort studies and bias the observed association toward the null.<sup>49 50</sup> Therefore, the true association between coffee consumption and risk of prostate cancer may be even stronger. Fourth, since coffee intake and the incidence of prostate cancer in the USA and Europe are relatively high, most of the studies were conducted in these regions. Since the effect size is small, we should be cautious when generalising the results to other areas, especially where the incidence is relatively low. Last, significant between-study heterogeneity may limit interpretation of the results. The observed heterogeneity may come from various sources. For example, the highest and lowest categories of coffee intake are different in the original studies. Studies with a broader range between the highest and lowest categories were assumed to generate a higher risk estimate. Moreover, the type of coffee and different brewing methods included in the coffee consumption groups differed. The different cohort sizes and follow-up periods may also lead to heterogeneous results. Taken together, due to the significant heterogeneity in the current meta-analysis, the pooled results should be interpreted with caution.

## CONCLUSIONS

This study suggests that increased coffee consumption may be associated with a reduced risk of prostate cancer. Further research is still warranted to explore the underlying mechanisms and active compounds in coffee. If the association is further proved to be a causal effect, men might be encouraged to increase their coffee consumption to potentially decrease the risk of prostate cancer.

**Correction notice** This article has been corrected since it first published. The provenance and peer review statement has been included.

**Acknowledgements** We would like to thank the authors of the original studies for their contribution to our meta-analysis, especially those authors who provided their raw data for the analysis.

**Contributors** KW obtained the funding, developed the research design, interpreted the results and also had primary responsibility for the final content. XC, YZ and ZT analysed the data and interpreted the results. XC and KW drafted manuscript. All authors critically reviewed and approved the manuscript.

**Funding** This work was supported by the Natural Science Foundation of Liaoning Province of China (Grant No 2019-MS-3608) for Kefeng Wang and 345 Talent Project of Shengjing Hospital of China Medical University for Kefeng Wang (Grant No M0122).

**Competing interests** None declared.

**Patient consent for publication** Not required.

**Provenance and peer review** Not commissioned; externally peer reviewed.

**Data availability statement** Data are available upon reasonable request. The data are available upon request from the corresponding author (wangkefenguro@sina.com)

**Supplemental material** This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

**Open access** This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>.

## ORCID iD

Kefeng Wang <http://orcid.org/0000-0003-3134-356X>

## REFERENCES

- Bray F, Ferlay J, Soerjomataram I, et al. Global cancer statistics 2018: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA Cancer J Clin* 2018;68:394–424.
- Cullen J, Elsamouni S, Brassell SA, et al. The burden of prostate cancer in Asian nations. *J Carcinog* 2012;11:7.
- Tunncliffe JM, Shearer J. Coffee, glucose homeostasis, and insulin resistance: physiological mechanisms and mediators. *Appl Physiol Nutr Metab* 2008;33:1290–300.
- Lee WJ, Zhu BT. Inhibition of DNA methylation by caffeic acid and chlorogenic acid, two common catechol-containing coffee polyphenols. *Carcinogenesis* 2006;27:269–77.
- Bode AM, Dong Z. The enigmatic effects of caffeine in cell cycle and cancer. *Cancer Lett* 2007;247:26–39.
- Yu X, Bao Z, Zou J, et al. Coffee consumption and risk of cancers: a meta-analysis of cohort studies. *BMC Cancer* 2011;11:96.
- Jacobsen BK, Bjelke E, Kvåle G, et al. Coffee drinking, mortality, and cancer incidence: results from a Norwegian prospective study. *J Natl Cancer Inst* 1986;76:823–31.



- 8 Nomura A, Heilbrun LK, Stemmermann GN. Prospective study of coffee consumption and the risk of cancer. *J Natl Cancer Inst* 1986;76:587–90.
- 9 Severson RK, Nomura AM, Grove JS, *et al.* A prospective study of demographics, diet, and prostate cancer among men of Japanese ancestry in Hawaii. *Cancer Res* 1989;49:1857–60.
- 10 Hsing AW, McLaughlin JK, Schuman LM, *et al.* Diet, tobacco use, and fatal prostate cancer: results from the Lutheran brotherhood cohort study. *Cancer Res* 1990;50:6836–40.
- 11 Le Marchand L, Kolonel LN, Wilkens LR, *et al.* Animal fat consumption and prostate cancer: a prospective study in Hawaii. *Epidemiology* 1994;5:276–82.
- 12 Stensvold I, Jacobsen BK. Coffee and cancer: a prospective study of 43,000 Norwegian men and women. *Cancer Causes Control* 1994;5:401–8.
- 13 Ellison LF. Tea and other beverage consumption and prostate cancer risk: a Canadian retrospective cohort study. *Eur J Cancer Prev* 2000;9:125–30.
- 14 Iso H, Kubota Y, Japan Collaborative Cohort Study for Evaluation of Cancer. Nutrition and disease in the Japan Collaborative Cohort Study for Evaluation of Cancer (JACC). *Asian Pac J Cancer Prev* 2007;8:35–80.
- 15 Nilsson LM, Johansson I, Lenner P, *et al.* Consumption of filtered and boiled coffee and the risk of incident cancer: a prospective cohort study. *Cancer Causes Control* 2010;21:1533–44.
- 16 Bosire C, Stampfer MJ, Subar AF, *et al.* Coffee consumption and the risk of overall and fatal prostate cancer in the NIH-AARP Diet and Health study. *Cancer Causes Control* 2013;24:1527–34.
- 17 Discacciati A, Orsini N, Andersson S-O, *et al.* Coffee consumption and risk of localized, advanced and fatal prostate cancer: a population-based prospective study. *Ann Oncol* 2013;24:1912–8.
- 18 Wilson KM, Kasperzyk JL, Rider JR, *et al.* Coffee consumption and prostate cancer risk and progression in the Health Professionals Follow-up Study. *J Natl Cancer Inst* 2011;103:876–84.
- 19 Shafique K, McLoone P, Qureshi K, *et al.* Coffee consumption and prostate cancer risk: further evidence for inverse relationship. *Nutr J* 2012;11:42.
- 20 Russnes KM, Wilson KM, Epstein MM, *et al.* Total antioxidant intake in relation to prostate cancer incidence in the Health Professionals Follow-up Study. *Int J Cancer* 2014;134:1156–65.
- 21 Li Q, Kakizaki M, Sugawara Y, *et al.* Coffee consumption and the risk of prostate cancer: the Ohsaki cohort study. *Br J Cancer* 2013;108:2381–9.
- 22 Park C-H, Myung S-K, Kim T-Y, *et al.* Coffee consumption and risk of prostate cancer: a meta-analysis of epidemiological studies. *BJU Int* 2010;106:762–9.
- 23 Liu H, Hu G-H, Wang X-C, *et al.* Coffee consumption and prostate cancer risk: a meta-analysis of cohort studies. *Nutr Cancer* 2015;67:392–400.
- 24 Tverdal A. Boiled coffee consumption and the risk of prostate cancer: follow-up of 224,234 Norwegian men 20–69 years. *Br J Cancer* 2015;112:576–9.
- 25 Hashibe M, Galeone C, Buys SS, *et al.* Coffee, tea, caffeine intake, and the risk of cancer in the PLCO cohort. *Br J Cancer* 2015;113:809–16.
- 26 Pounis G, Tabolacci C, Costanzo S, *et al.* Reduction by coffee consumption of prostate cancer risk: evidence from the Moli-sani cohort and cellular models. *Int J Cancer* 2017;141:72–82.
- 27 Sen A, Papadimitriou N, Lagiou P, *et al.* Coffee and tea consumption and risk of prostate cancer in the European Prospective Investigation into Cancer and Nutrition. *Int J Cancer* 2019;144:240–50.
- 28 Ong J-S, Law MH, An J, *et al.* Association between coffee consumption and overall risk of being diagnosed with or dying from cancer among >300 000 UK Biobank participants in a large-scale Mendelian randomization study. *Int J Epidemiol* 2019;48:1447–56.
- 29 Moher D, Liberati A, Tetzlaff J, *et al.* Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLoS Med* 2009;6:e1000097.
- 30 Wells GA, Shea B, O'Connell D. The Newcastle-Ottawa scale (NOS) for assessing the quality of nonrandomised studies in meta-analyses [http://www.ohri.ca/programs/clinical\\_epidemiology/oxford.asp](http://www.ohri.ca/programs/clinical_epidemiology/oxford.asp)
- 31 DerSimonian R, Laird N. Meta-analysis in clinical trials. *Control Clin Trials* 1986;7:177–88.
- 32 Welch HG, Albertsen PC. Prostate cancer diagnosis and treatment after the introduction of prostate-specific antigen screening: 1986–2005. *J Natl Cancer Inst* 2009;101:1325–9.
- 33 Crippa A, Discacciati A, Larsson SC, *et al.* Coffee consumption and mortality from all causes, cardiovascular disease, and cancer: a dose-response meta-analysis. *Am J Epidemiol* 2014;180:763–75.
- 34 Kennedy OJ, Roderick P, Buchanan R, *et al.* Coffee, including caffeinated and decaffeinated coffee, and the risk of hepatocellular carcinoma: a systematic review and dose-response meta-analysis. *BMJ Open* 2017;7:e013739.
- 35 Greenland S, Longnecker MP. Methods for trend estimation from summarized dose-response data, with applications to meta-analysis. *Am J Epidemiol* 1992;135:1301–9.
- 36 Orsini N, Li R, Wolk A, *et al.* Meta-analysis for linear and nonlinear dose-response relations: examples, an evaluation of approximations, and software. *Am J Epidemiol* 2012;175:66–73.
- 37 Higgins JPT, Thompson SG. Quantifying heterogeneity in a meta-analysis. *Stat Med* 2002;21:1539–58.
- 38 Begg CB, Mazumdar M. Operating characteristics of a RANK correlation test for publication bias. *Biometrics* 1994;50:1088–101.
- 39 Egger M, Davey Smith G, Schneider M, *et al.* Bias in meta-analysis detected by a simple, graphical test. *BMJ* 1997;315:629–34.
- 40 Endogenous Hormones and Prostate Cancer Collaborative Group, Roddam AW, Allen NE, *et al.* Endogenous sex hormones and prostate cancer: a collaborative analysis of 18 prospective studies. *J Natl Cancer Inst* 2008;100:170–83.
- 41 Williams CJ, Fargnoli JL, Hwang JJ, *et al.* Coffee consumption is associated with higher plasma adiponectin concentrations in women with or without type 2 diabetes: a prospective cohort study. *Diabetes Care* 2008;31:504–7.
- 42 Imatoh T, Tanihara S, Miyazaki M, *et al.* Coffee consumption but not green tea consumption is associated with adiponectin levels in Japanese males. *Eur J Nutr* 2011;50:279–84.
- 43 Ziemke F, Mantzoros CS. Adiponectin in insulin resistance: lessons from translational research. *Am J Clin Nutr* 2010;91:258S–61.
- 44 Ma J, Li H, Giovannucci E, *et al.* Prediagnostic body-mass index, plasma C-peptide concentration, and prostate cancer-specific mortality in men with prostate cancer: a long-term survival analysis. *Lancet Oncol* 2008;9:1039–47.
- 45 Hammarsten J, Högstäd B. Hyperinsulinaemia: a prospective risk factor for lethal clinical prostate cancer. *Eur J Cancer* 2005;41:2887–95.
- 46 Thapa D, Ghosh R. Antioxidants for prostate cancer chemoprevention: challenges and opportunities. *Biochem Pharmacol* 2012;83:1319–30.
- 47 Svartberg J, Midtby M, Bønna KH, *et al.* The associations of age, lifestyle factors and chronic disease with testosterone in men: the Tromsø study. *Eur J Endocrinol* 2003;149:145–52.
- 48 Goto A, Song Y, Chen BH, *et al.* Coffee and caffeine consumption in relation to sex hormone-binding globulin and risk of type 2 diabetes in postmenopausal women. *Diabetes* 2011;60:269–75.
- 49 Qi X-X, Shen P. Associations of dietary protein intake with all-cause, cardiovascular disease, and cancer mortality: a systematic review and meta-analysis of cohort studies. *Nutr Metab Cardiovasc Dis* 2020;30:1094–105.
- 50 Yang Y, Zhao L-G, Wu Q-J, *et al.* Association between dietary fiber and lower risk of all-cause mortality: a meta-analysis of cohort studies. *Am J Epidemiol* 2015;181:83–91.



## PRISMA 2009 Checklist

Section/topic	#	Checklist item	Reported on page #
<b>TITLE</b>			
Title	1	Identify the report as a systematic review, meta-analysis, or both.	P1, L1-2
<b>ABSTRACT</b>			
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.	P2, 3, L20-46
<b>INTRODUCTION</b>			
Rationale	3	Describe the rationale for the review in the context of what is already known.	P5, L84-87
Objectives	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).	P5, L88-91
<b>METHODS</b>			
Protocol and registration	5	Indicate if a review protocol exists, if and where it can be accessed (e.g., Web address), and, if available, provide registration information including registration number.	NA
Eligibility criteria	6	Specify study characteristics (e.g., PICOS, length of follow-up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rationale.	P6, L104-125
Information sources	7	Describe all information sources (e.g., databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched.	P7, L127-140
Search	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	P7, L127-13
Study selection	9	State the process for selecting studies (i.e., screening, eligibility, included in systematic review, and, if applicable, included in the meta-analysis).	P6, L105-124
Data collection process	10	Describe method of data extraction from reports (e.g., piloted forms, independently, in duplicate) and any processes for obtaining and confirming data from investigators.	P5, L97-98
Data items	11	List and define all variables for which data were sought (e.g., PICOS, funding sources) and any assumptions and simplifications made.	P7, 8, L142-148
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.	P8, L150-153
Summary measures	13	State the principal summary measures (e.g., risk ratio, difference in means).	P8,



## PRISMA 2009 Checklist

			L155-156
Synthesis of results	14	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., $I^2$ ) for each meta-analysis.	P8, 9, L156-188

Page 1 of 2

Section/topic	#	Checklist item	Reported on page #
Risk of bias across studies	15	Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies).	P9, L184-185
Additional analyses	16	Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified.	P9, L170-182
<b>RESULTS</b>			
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.	P10, L192-204
Study characteristics	18	For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citations.	P10, 11, L206-216
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).	P11, L215-216
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.	P11, L218-219
Synthesis of results	21	Present results of each meta-analysis done, including confidence intervals and measures of consistency.	P11, L220-227
Risk of bias across studies	22	Present results of any assessment of risk of bias across studies (see Item 15).	P11, L227-230
Additional analysis	23	Give results of additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression [see Item 16]).	P11, 12, L232-244
<b>DISCUSSION</b>			
Summary of evidence	24	Summarize the main findings including the strength of evidence for each main outcome; consider their relevance to key groups (e.g., healthcare providers, users, and policy makers).	P14, L259-263
Limitations	25	Discuss limitations at study and outcome level (e.g., risk of bias), and at review-level (e.g., incomplete retrieval of identified research, reporting bias).	P17, 18, L322-354
Conclusions	26	Provide a general interpretation of the results in the context of other evidence, and implications for future research.	P18,



## PRISMA 2009 Checklist

			L357-361
<b>FUNDING</b>			
Funding	27	Describe sources of funding for the systematic review and other support (e.g., supply of data); role of funders for the systematic review.	P19, L364-366

From: 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.pmed1000097

For more information, visit: [www.prisma-statement.org](http://www.prisma-statement.org).

Page 2 of 2



## MOOSE Guidelines for Meta-Analyses and Systematic Reviews of Observational Studies\*

	Topic	P1, L1-2 Page number
<b>Title</b>	Identify the study as a meta-analysis (or systematic review)	P2-3, L20-40
<b>Abstract</b>	Use the journal's structured format	P5, L84-87
<b>Introduction</b>	<b>Present:</b>	P5, L87-88
	The clinical problem	P5, L88-90
	The hypothesis	P5, L88-90
	A statement of objectives that includes the study population, the condition of interest, the exposure or intervention, and the outcome(s) considered	
<b>Sources</b>	<b>Describe:</b>	P5, L96
	Qualifications of searchers (eg, librarians and investigators)	P7, L129-130
	Search strategy, including time period included in the synthesis and keywords	P7, L133-140
	Effort to include all available studies, including contact with authors	P7, L127-128
	Databases and registries searched	
	Search software used, name and version, including special features used (e.g. explosion)	P7, L128-130
	Use of hand searching (e.g. reference lists of obtained articles)	P7, L130-131
	List of citations located and those excluded, including justification	NA
	Method of addressing articles published in languages other than English	NA
	Method of handling abstracts and unpublished studies	P7, L133-134
	Description of any contact with authors	P7, L133-140
<b>Study Selection</b>	<b>Describe</b>	
	Types of study designs considered	P6, L105-106
	Relevance or appropriateness of studies gathered for assessing the hypothesis to be tested	P6, L105-124
	Rationale for the selection and coding of data (eg, sound clinical principles or convenience)	NA
	Documentation of how data were classified and coded (eg, multiple raters, blinding, and inter-rater reliability)	P5, L96-99
	Assessment of confounding (e.g. comparability of cases and controls in studies where appropriate)	P8, L161-167
	Assessment of study quality, including blinding of quality assessors; stratification or regression on possible predictors of study results	P8, L149-153
	Assessment of heterogeneity	
	Statistical methods (eg, complete description of fixed or random effects models, justification of whether the chosen models account for predictors of study results, dose-response models, or cumulative meta-analysis) in sufficient detail to be replicated	P9, L183-184
<b>Results</b>	<b>Present</b>	P8-9, L155-188
	A graph summarizing individual study estimates and the overall estimate	
	A table giving descriptive information for each included study	
	Results of sensitivity testing (eg, subgroup analysis)	Figure 2
	Indication of statistical uncertainty of findings	Supplementary Document 4
<b>Discussion</b>	<b>Discuss</b>	P11-12, L231-244
	Strengths and weaknesses	P11-12, L217-244
	Potential biases in the review process (eg, publication bias)	

P16-18, L312-353

P17, L324-341

P17, L320-321

P18, L344-346

	Assessment of quality of included studies	P18, L346-354
	Consideration of alternative explanations for observed results	
	Generalization of the conclusions (ie, appropriate for the data presented and within the domain of the literature review)	P18, L358-361
	Guidelines for future research	P19, L364-366
	Disclosure of funding source	

\*Modified from Stroup DF, Berlin JA, Morton SC, Olkin I, Williamson GD, Rennie D, et al. Meta-analysis of observational studies in epidemiology: a proposal for reporting. Meta-analysis Of Observational Studies in Epidemiology (MOOSE) group. JAMA 2000;283:2008–12. Copyrighted © 2000, American Medical Association. All rights reserved.

**Pubmed (N=116)****Search date:** up to Sep 21, 2020**Search terms:**

("coffee"[MeSH Terms] OR "coffee"[All Fields] OR "coffee s"[All Fields] OR "coffees"[All Fields]) AND ("prostat"[All Fields] OR "prostate"[MeSH Terms] OR "prostate"[All Fields] OR "prostates"[All Fields] OR "prostatic"[All Fields] OR "prostatism"[MeSH Terms] OR "prostatism"[All Fields] OR "prostatitis"[MeSH Terms] OR "prostatitis"[All Fields]) AND ("cancer s"[All Fields] OR "cancerated"[All Fields] OR "canceration"[All Fields] OR "cancerization"[All Fields] OR "cancerized"[All Fields] OR "cancerous"[All Fields] OR "neoplasms"[MeSH Terms] OR "neoplasms"[All Fields] OR "cancer"[All Fields] OR "cancers"[All Fields] OR ("carcinoma"[MeSH Terms] OR "carcinoma"[All Fields] OR "carcinomas"[All Fields] OR "carcinoma s"[All Fields]) OR ("neoplasm s"[All Fields] OR "neoplasms"[MeSH Terms] OR "neoplasms"[All Fields] OR "neoplasm"[All Fields]) OR ("cysts"[MeSH Terms] OR "cysts"[All Fields] OR "cyst"[All Fields] OR "neurofibroma"[MeSH Terms] OR "neurofibroma"[All Fields] OR "neurofibromas"[All Fields] OR "tumor s"[All Fields] OR "tumoral"[All Fields] OR "tumorous"[All Fields] OR "tumour"[All Fields] OR "neoplasms"[MeSH Terms] OR "neoplasms"[All Fields] OR "tumor"[All Fields] OR "tumour s"[All Fields] OR "tumoural"[All Fields] OR "tumorous"[All Fields] OR "tumours"[All Fields] OR "tumors"[All Fields]))

**Web of Science (N=162)****Search date:** up to Sep 21, 2020**Search terms:**

TOPIC: (coffee) AND TOPIC: (prostate) AND TOPIC: (cancer OR carcinoma OR neoplasm OR tumor) Indexes=SCI-EXPANDED, CPCI-S, CCR-EXPANDED, IC Timespan=All years

**Embase (N=219)****Search date:** up to Sep 21, 2020**Search terms:**

coffee AND prostate AND (cancer OR carcinoma OR neoplasm OR tumor)

Table 1. Characteristics of cohort studies of coffee consumption and prostate cancer risk included in the meta-analysis

Study	Country	Study period	Age at baseline	Size of cohort/controls	No. of cases	Exposure assessment methods	Definition of coffee consumption	Outcome	Follow-up time	Confounder adjustments	NOS total quality score; Risk of bias (Potential bias)
Ong et al. 2019	UK	2006-2010	37-73 years	131834	7532	Self-reported diet survey	1 cup/day increase; no information on the highest and lowest coffee intakes	Total prostate cancer	<5 years	Age, townsend deprivation index, top 10 ancestral principal components, smoking status, BMI, height, alcohol intake, drink temperature, overall heath rating, highest qualification. Instrumental variable analyses (SNP instruments) were also used to control confounders	7 stars; low risk of bias (exposure misclassification bias)
Sen et al. 2019	Europe	1990s-2015	Mean: 52 years.	142196	7036	Validated FFQ	The highest intake: median of 855 ml/day (no. of cases: 1271); The lowest intake: median of 0 ml/day (no. of cases: 396)	Total, Localized, advanced prostate cancer	Mean: 14 years	Stratified by center and age at recruitment in 5 years categories, and adjusted for smoking status, BMI, history of diabetes, alcohol intake, education, physical activity, energy intake, as well as calcium, fish, tea, fruit and vegetable intake.	9 stars; low risk of bias
Pounis et al 2017	Italy	2005-2010	≥50 years;	6989	100	Validated FFQ	The highest intake: >3 cups/day (>90	Total prostate cancer	Mean: 4.24 years	Age, energy intake, smoking habits and BMI	8 stars; low risk of bias



			Mean: 67 years				g/day) (no. of cases: 14); The lowest intake: 0-2 cups/day (0- 55 g/day) (no. of cases: 45)				
Hashibe et al. 2015	USA	1992-2011	55-74 years	46771	3037	Validated diet history questionnaire	Mean coffee intake is 1.9 cups/day; The highest intake: ≥2 cups/day (no. of cases: 1731); The lowest intake: <1 cups/day (no. of cases: 889)	Total prostate cancer	>10 years	Age, sex, race, and education.	8 stars; low risk of bias (confounding bias)
Tverdal et al. 2015	Norway	1974-1999	20-69 years	224234	5740	Questionnaire	The highest intake: ≥9 cups/day (no. of cases: 642); The lowest intake: none (no. of cases: 389)	Total prostate cancer	Mean: 17.6 years	Age, smoking, BMI, height, physical activity, total cholesterol, triglycerides, systolic blood pressure, year of examination and diabetes	8 stars; low risk of bias (exposure misclassification bias)
Li et al. 2013	Japan	1995-2005	40-79 years	18,853	318	Validated FFQ	The highest intake: ≥3	Total prostate	11 years	Age, education, BMI, time engaging in sports or exercise, marital status, time spent walking,	8 stars; low risk of bias

							cups/day (no. of cases: 24); The lowest intake: none (no. of cases: 84)	cancer incidence			smoking status, family history of cancer, job status, total energy intake, passive smoking, alcohol drinking, daily consumption of miso soup	
Discacciati et al. 2013	Sweden	1998-2010	45-79 years	44,613	3801	Validated self-administered FFQ	The highest intake: ≥6 cups/day (median of 1484 g/day) (no. of cases: 173); The lowest intake: none (median: 0 g/day) (no. of cases: 129)	Localized and advanced prostate cancer incidence	13 years	Age, tea, alcohol, BMI, diabetes, family history of prostate cancer, smoke, physical activity, education, total energy intake.	8 stars; low risk of bias	
Bosire et al. 2013	USA	1995-2008	50-71 years	288,391	23335	Validated FFQ	The highest intake: ≥6 cups/day (no. of cases: 787); The lowest intake: none (no. of cases: 2136)	Total prostate cancer incidence	>11 years (median: 10.5 years)	Age, race, height, BMI, physical activity, smoking, history of diabetes, family history of prostate cancer, PSA testing, intakes of tomato sauce, alpha-linolenic acid, and total energy intake.	8 stars; low risk of bias	
Shafique et al. 2012	UK	1970-2007	21-75 years (median	6017	318	Self-administered questionnaire	The highest intake: ≥3 cups/day (no. of cases: 65);	Total prostate cancer incidence	37 years (median: 28 years)	Age at screening, cholesterol, systolic blood pressure, BMI, alcohol intake, tea consumption, smoking status, social class.	8 stars; low risk of bias (exposure misclassification bias)	

			: 48 years)				The lowest intake: none (no. of cases: 139)				
Wilson et al. 2011	USA	1986-2006	40-75 years	47,911	5035	Validated FFQ	The highest intake: ≥6 cups/day (no. of cases: 152); The lowest intake: none (no. of cases: 587)	Total prostate cancer incidence	20 years	Age in months, calendar time, race, BMI at age 21, current BMI, vigorous physical activity, smoking, diabetes, family history of prostate cancer in father or brother, multivitamin use, intakes of processed meat, tomato sauce, calcium, alpha-linolenic acid, supplemental vitamin E, alcohol intake, energy intake, history of PSA testing.	9 stars; low risk of bias
Nilsson et al. 2010	Sweden	1985-2007	40-60 years (median : 50 years)	30,930	653	Validated Semi- quantitative FFQ	The highest intake: ≥4 cups/day (no. of cases: 209); The lowest intake: <1 cup/day (no. of cases: 60)	Total prostate cancer incidence	15 years (median: 6 years)	Age, BMI, smoking, education, recreational physical activity.	8 stars; low risk of bias
Iso et al. 2007	Japan	1988-1997	40-79 years	43,500	161	Self- administrated questionnaire	The highest intake: ≥2 cups/day (no. of cases: 38);	Prostate cancer mortality	Mean: 8.15 years	Age, area of study	7 stars; low risk of bias (exposure misclassification bias, confounding bias)

The lowest intake: ≤1-2 cup/month (no. of cases: 47)											
Ellison et al. 2000	Canada	1970-1993	50-84 years	3400	145	FFQ	The highest intake: ≥750 mg/day (no. of cases: 122); The lowest intake: 0 mg/day (no. of cases: 23)	Total prostate cancer incidence	Mean: 11.6 year	Age, wine consumption.	6 stars; medium risk of bias (exposure misclassification bias, confounding bias)
Le Marchand et al. 1994	USA	1975-1989	≥45 years	20,316	198	Self- administered life-style questionnaire	The highest intake: ≥2.5 cups/day; The lowest intake: none.	Total prostate cancer incidence	Median: 6 years	Age, ethnicity, income.	6 stars; medium risk of bias (exposure misclassification bias, confounding bias)
Hsing et al. 1990	USA	1966-1986	≥35 years (Median : 51 years)	17,633	149	FFQ	The highest intake: ≥5 cups/day; The lowest intake: <3 cups/day.	Prostate cancer mortality	20 years (Mean: 15.6 years)	Age, tobacco use.	7 stars; low risk of bias (exposure misclassification bias, confounding bias)
Severson et al. 1989	USA	1965-1986	46-68 years	7998	174	FFQ + 24-h diet recall interview	The highest intake: ≥5 cups/week (no. of cases: 146);	Total prostate cancer incidence	Mean: 17.4 years	Age	7 stars; low risk of bias (confounding bias)



The lowest intake:

<1 cups/week (no.

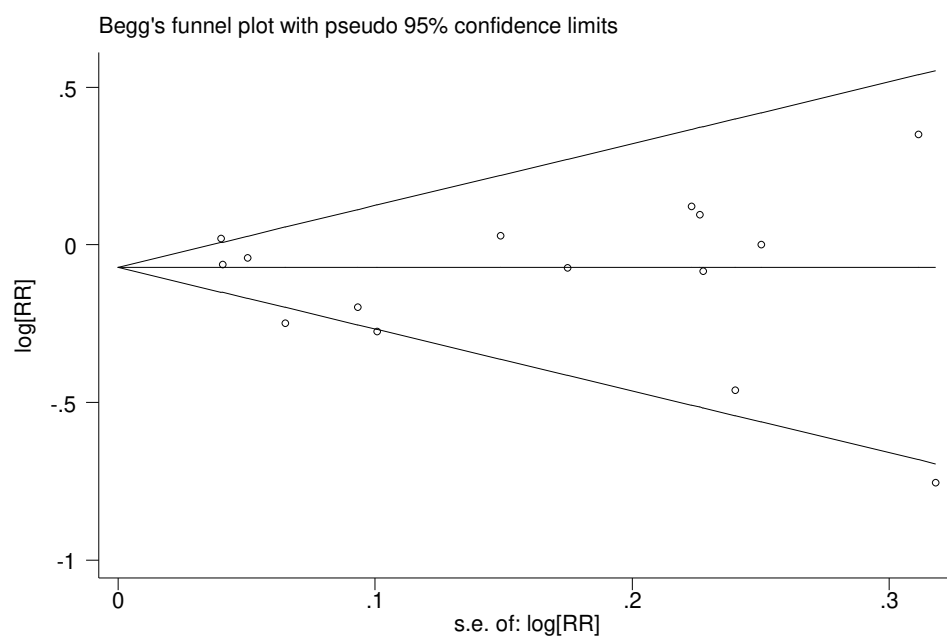
of cases: 22)

BMI, body mass index; CI, confidence interval; FFQ, food frequency questionnaire; NA, not available; PSA, prostate-specific antigen; RR, relative risk;

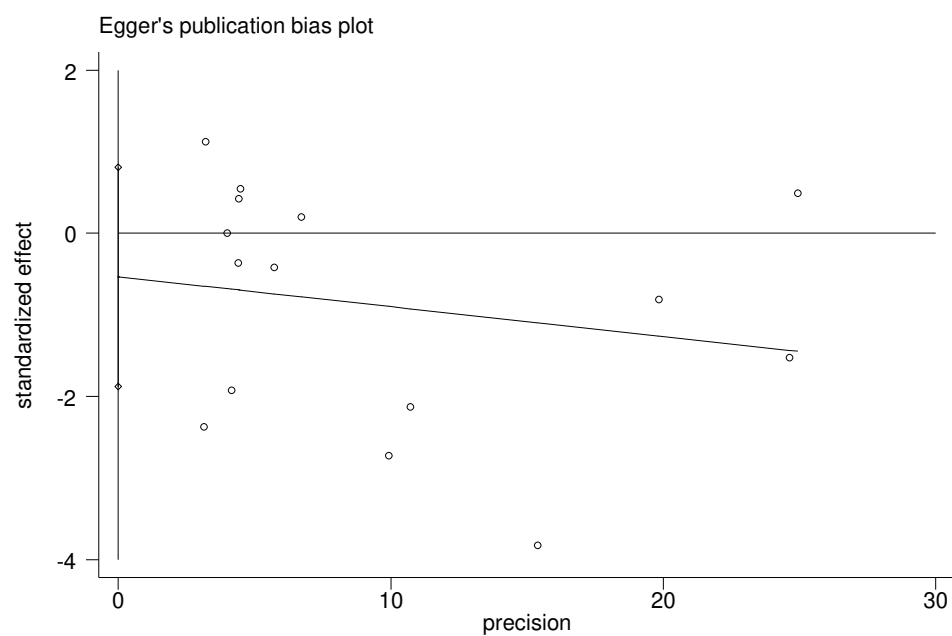
Supplementary Table S3 Quality of cohort studies included in the meta-analysis

Study	Representativeness of the exposed cohort	Selection of the non-exposed cohort	Ascertainment of exposure	Outcome of interest was not present at start of study	Controls for important risk factors <sup>1</sup>	Assessment of outcome	Follow-up long enough for outcomes to occur	Adequacy of follow up of cohorts	Total quality score
Ong et al. 2019	☆	☆	–	☆	☆☆	☆	–	☆	7
Sen et al. 2019	☆	☆	☆	☆	☆☆	☆	☆	☆	9
Pounis et al 2017	☆	☆	☆	☆	☆☆	☆	–	☆	8
Hashibe et al. 2015	☆	☆	☆	☆	☆	☆	☆	☆	8
Terdal et al. 2015	☆	☆	–	☆	☆☆	☆	☆	☆	8
Li et al. 2013	☆	☆	☆	☆	☆☆	☆	☆	–	8
Discacciati et al. 2013	☆	☆	☆	☆	☆☆	☆	☆	–	8
Bosire et al. 2013	☆	☆	☆	☆	☆☆	☆	☆	–	8
Shafique et al. 2012	☆	☆	–	☆	☆☆	☆	☆	☆	8
Wilson et al. 2011	☆	☆	☆	☆	☆☆	☆	☆	☆	9
Nilsson et al. 2010	☆	☆	☆	–	☆☆	☆	☆	☆	8
Iso et al. 2007	☆	☆	–	☆	☆	☆	☆	☆	7
Ellison et al. 2000	☆	☆	–	☆	☆	☆	☆	–	6
Le Marchand et al. 1994	☆	☆	–	☆	☆	☆	☆	–	6
Hsing et al. 1990	☆	☆	–	☆	☆	☆	☆	☆	7
Severson et al. 1989	☆	☆	☆	☆	☆	☆	☆	–	7

1. A maximum of 2 stars could be awarded for this item. Studies that included adjustment for age received one star, and studies that included most of the other important confounders such as ethnicity, dietary factors (energy intake, vitamin D, dietary fat etc.), physical activity, body mass index, type 2 diabetes mellitus, alcohol and smoking received an additional star.



**Figure S1** Begg's funnel plot of coffee consumption and prostate cancer risk



**Figure S2** Egger's publication bias plot of coffee consumption and prostate cancer risk