Mediterranean Diet Score and prostate cancer risk in a Swedish population-based case–control study

Elisabeth Möller1*, Carlotta Galeone2,3, Therese M.-L. Andersson1, Rino Bellocco1,4, Hans-Olov Adami1,5, Ove Andrén6, Henrik Grönberg1, Carlo La Vecchia2,3, Lorelei A. Mucci5,7 and Katarina Bälter1

1Department of Medical Epidemiology and Biostatistics, Karolinska Institutet, PO Box 281, Stockholm SE-171 77, Sweden
2Department of Epidemiology, Mario Negri Institute for Pharmacological Research, Milan, Italy
3Department of Clinical Sciences and Community Health, University of Milan, Milan, Italy
4Department of Statistics and Quantitative Methods, University of Milano-Bicocca, Milan, Italy
5Department of Epidemiology, Harvard School of Public Health, Boston, MA, USA
6Department of Urology, Örebro University Hospital, Örebro, Sweden
7Department of Medicine, Channing Laboratory, Brigham and Women’s Hospital and Harvard Medical School, Boston, MA, USA

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Abstract
Several individual components of the Mediterranean diet have been shown to offer protection against prostate cancer. The present study is the first to investigate the association between adherence to the Mediterranean diet and the relative risk of prostate cancer. We also explored the usefulness of the Mediterranean Diet Score (MDS) in a non-Mediterranean population. FFQ data were obtained from 1482 incident prostate cancer patients and 1108 population-based controls in the Cancer of the Prostate in Sweden (CAPS) study. We defined five MDS variants with different components or using either study-specific intakes or intakes in a Greek reference population as cut-off values between low and high intake of each component. Unconditional logistic regression was used to estimate the relative risk of prostate cancer for high and medium vs. low MDS, as well as potential associations with the individual score components. No statistically significant association was found between adherence to the Mediterranean diet based on any of the MDS variants and prostate cancer risk (OR range: 0.96–1.19 for total prostate cancer, comparing high with low adherence). Overall, we found little support for an association between the Mediterranean diet and prostate cancer in this Northern European study population. Despite potential limitations inherent in the study or in the build-up of a dietary score, we suggest that the original MDS with study-specific median intakes as cut-off values between low and high intake is useful in assessing the adherence to the Mediterranean diet in non-Mediterranean populations.

Key words: Prostatic neoplasms; Dietary patterns; Dietary score; Epidemiology

Prostate cancer is the second most common cancer among men globally and is the most common cancer in high-income countries(1). However, its incidence varies considerably in different parts of the world. The highest incidence is seen in the Nordic countries, North America, Australia and New Zealand, with age-standardised rates ranging from seventy-three to 112 per 100,000 person-years(2). In most Mediterranean countries, the incidence rates are almost half to two-thirds lower(3), suggesting that certain aspects of the Mediterranean lifestyle may be associated with a reduced incidence of prostate cancer.

A number of epidemiological studies suggest that a Mediterranean dietary pattern may offer protection against several types of cancer(2,3). The traditional Mediterranean diet is the dietary habit typical of the Mediterranean regions, notably

Abbreviations: CAPS, Cancer of the Prostate in Sweden; MDS, Mediterranean Diet Score; MPS, MUFA and PUFA to SFA.

* Corresponding author: Elisabeth Möller, fax + 46 8 31 49 75, email elisabeth.moller@ki.se

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Crete and other parts of Greece, southern Italy and Spain, in the period before the 1960s. The general features of this diet are high intake of vegetables, fruits, nuts, beans, cereals and lean fish, together with high consumption of olive oil, moderate consumption of alcohol (especially wine with meals) and low amounts of milk and red meat. Several of these individual food items, or the nutrients they contain, have been associated with a reduced risk of prostate cancer. Based on previous knowledge about the effect of Mediterranean dietary components, Trichopoulou et al. estimated that shifting to a traditional Mediterranean diet could prevent up to 10% of the prostate cancer cases in Western high-income countries.

Several dietary scores have been developed to assess adherence to the Mediterranean diet. In 1995, Trichopoulou et al. developed the Mediterranean Diet Score (MDS) based on eight dietary components: the intake of vegetables, fruits and nuts, legumes, cereals, meat, dairy products and alcohol, as well as the ratio of MUFA to SFA. Fish was added as a ninth component in 2003. The MDS and several modified versions have been used in epidemiological studies of various health outcomes, both in Mediterranean and non-Mediterranean populations. An important component of the traditional Mediterranean diet is olive oil, resulting in a high ratio of MUFA to SFA. Outside the Mediterranean region olive oil is consumed less frequently, and the intake of MUFA is not only considerably lower, but mainly originates from a higher meat intake rather than from vegetable oils. In an attempt to adapt the MDS for use in non-Mediterranean populations, the fat ratio component has been modified so as to include both MUFA and PUFA in the numerator and SFA in the denominator. Moreover, the cut-off value between high and low intake for each MDS component is the median intake in the population under study. However, using the median intake in a non-Mediterranean study population may be questionable, as the intake of certain components may be much lower or higher than that in a typical Mediterranean population. Thus the score may be less able to discriminate between intake levels that are beneficial or non-beneficial to health.

In the present study, we investigated the association between the Mediterranean diet and the relative risk of prostate cancer in a Swedish population. In order to examine the potential effect of using study-specific median intakes in a non-Mediterranean population as cut-off values in the MDS, we developed several variants of the score using both intakes in the Swedish study population and in a Greek reference population as cut-off values. We also created an alternative score that more closely reflects the traditional Mediterranean diet, directed towards its most beneficial components. Furthermore, we explored the potential effect measure modification by selected covariates. To the best of our knowledge, this is the first study specifically investigating prostate cancer risk and the Mediterranean diet using a dietary score.

Experimental methods

Study population

The Cancer of the Prostate in Sweden (CAPS) study is a population-based case–control study on prostate cancer and has been described in detail previously. Briefly, incident and histologically confirmed prostate cancer cases were identified from four of the six regional cancer registries in Sweden and were invited to take part in the study. Cases were 35–79 years old at enrolment, from January 2001 to September 2002, and lived in the central and northern parts of Sweden. For 95% of the cases, clinical data were obtained from the National Prostate Cancer Registry. Advanced cases were defined as those meeting at least one of the following criteria: tumour, nodes, metastasis stage T3/T4 or N1 or M1; Gleason score 8–10; or serum prostate-specific antigen level at diagnosis ≥100 ng/ml. Localised cases were defined as those with T1/T2, N0, M0, Gleason score 2–6 and prostate-specific antigen <20 ng/ml. Cases with Gleason score 7 who did not meet the criteria for advanced disease were included in neither advanced nor localised cases due to the difference in aggressiveness seen for Gleason 3 + 4 compared with Gleason 4 + 3. Controls were selected randomly from the Swedish Population Registry every 6 months, being frequency-matched to cases by age in 5-year categories and region of residence. Invitations to participate were sent out to eligible controls about once a month, except July and August, to reflect the continuous enrolment of cases.

All study participants were asked to complete a baseline questionnaire. The number of cases filling in the questionnaire was 1499 (79%) out of 1895 invited and the corresponding number for controls was 1130 (67%) out of 1684 invited. The average time between diagnosis and sending out the questionnaire was 5 months. The study was performed according to the guidelines laid down in the Declaration of Helsinki and was approved by the ethics committees at Karolinska Institutet and Umeå University in Sweden. Written informed consent was obtained from all participants.

Exposure assessment

Usual dietary intake over the past year was assessed using a semi-quantitative FFQ with 106 items, including foods, beverages and alcohol, and with three additional questions on dietary fat, including a question on regular use of olive oil for cooking and/or in dressings (yes/no). The intake of several food items assessed with a shorter version of the FFQ has been validated against fourteen repeated 24-h recall interviews in 248 randomly sampled Swedish women, showing Spearman correlation coefficients in the range 0.16–0.82 for vegetables/tomatoes, 0.38–0.49 for fruit/fruit juice, 0.16–0.61 for refined grains/whole grains/cereals, 0.44 for fish, 0.44 for dairy products, 0.37–0.60 for meat/poultry and 0.56–0.82 for alcoholic beverages. The intake of several food items assessed with a shorter version of the FFQ has been validated against four 7 d weighed food records in 111 randomly sampled Swedish women, giving Spearman correlation coefficients in the range 0.16–0.82 for vegetables/tomatoes, 0.38–0.49 for fruit/fruit juice, 0.16–0.61 for refined grains/whole grains/cereals, 0.44 for fish, 0.44 for dairy products, 0.37–0.60 for meat/poultry and 0.56–0.82 for alcoholic beverages. The intake of several food items assessed with a shorter version of the FFQ has been validated against four 7 d weighed food records in 111 randomly sampled Swedish women, giving Spearman correlation coefficients in the range 0.16–0.82 for vegetables/tomatoes, 0.38–0.49 for fruit/fruit juice, 0.16–0.61 for refined grains/whole grains/cereals, 0.44 for fish, 0.44 for dairy products, 0.37–0.60 for meat/poultry and 0.56–0.82 for alcoholic beverages. The intake of several food items assessed with a shorter version of the FFQ has been validated against four 7 d weighed food records in 111 randomly sampled Swedish women, giving Spearman correlation coefficients in the range 0.16–0.82 for vegetables/tomatoes, 0.38–0.49 for fruit/fruit juice, 0.16–0.61 for refined grains/whole grains/cereals, 0.44 for fish, 0.44 for dairy products, 0.37–0.60 for meat/poultry and 0.56–0.82 for alcoholic beverages.
We created five variants of the MDS originally developed by Trichopoulou et al. (24), as listed in Table 1. The main score, denoted as MDS-gram, as based on nine components: a high ratio of MUFA and PUFA to SFA (OMPS); high vegetable intake (including tomato juice, ketchup and root vegetables, except potatoes); high intake of fruit, nuts and seeds; high legume intake; high cereal intake; high fish and seafood intake; high intake of dairy products; high intake of meat and meat products (including poultry); and moderate alcohol intake. Frequencies of intake from the questionnaire were translated into scores of 0, 1, or 2, corresponding to zero, one, or two points for each component. One point was assigned to each component using the median intake among the controls. The total score was then categorised into low (zero to three points), medium (four to five points) or high (six to nine points) adherence to the Mediterranean diet. The MP:S ratio, ratio of MUFA and PUFA to SFA; CAPS, Cancer of the Prostate in Sweden study; serv/w, servings per week; alt, alternative.

Table 1. Description of the Mediterranean Diet Score (MDS) variants

| Score          | Unit          | Cut-off values | MP:S ratio | Vegetables* | Fruits | Nuts | Legumes | Cereals | Fish† | Dairy products | Meat/meat products‡ | Ethanol | Total score |
|----------------|---------------|----------------|------------|-------------|--------|------|---------|---------|-------|----------------|---------------------|---------|-------------|
| MDS-gram       | g/d           | Median in CAPS | x          | x           | +Nuts/seeds |        |       |         | x      | x               | x                   | x       | 0–9         |
| MDS-serv       | serv/w        | Median in CAPS | x          | x           | +Nuts/seeds |        |       |         | x      | x               | x                   | x       | 0–9         |
| MDS-cent       | g/d           | Twenty-fifth/  | x          | x           | +Nuts/seeds |        |       |         | x      | x               | x                   | x       | 0–9         |
| MDS-alt        | g/d           | Mean in Greek | x          | x           | +Nuts/seeds |        |       |         | x      | x               | x                   | x       | 0–9         |

† Includes vegetables, tomato juice, ketchup and root vegetables, except potatoes. Potatoes were not included among either vegetables or cereals, as consistent with the original score. However, including them in either of these groups did not change the results significantly.

‡ Includes cereal and seafood.

§ The twenty-fifth centile was used for the intake of legumes, cereals, fish, milk and meat, and the seventy-fifth centile for the MP:S ratio and the intake of vegetables, fruits/nuts and ethanol.

¶ The seventy-fifth centile was used as the cut-off value since the median intake was zero.

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centiles of the intake (in g/d) were used as cut-off values instead of the median intake; the centile level for each component was chosen so as to achieve cut-off values that were close to the median intakes in the Greek reference population, while maintaining a sufficient number of participants in the extreme groups. In MDS-greek, the median intakes (in g/d) in the aforementioned Greek reference population were used as cut-off values; however, for vegetable and fruit/nut intake we used the ninetieth centile of the intake in the CAPS population since very few participants reached the Greek median intake levels. MDS-serv and MDS-cent were categorised using the same adherence cut-off values as MDS-gram, whereas MDS-greek was categorised as low (zero to two points), medium (three to four points), or high (five to nine points) adherence.

Lastly, we created an alternative score, denoted as MDS-alt, with a focus on the most ‘traditional’ components of the Mediterranean diet. This score was based on ten components, basically the same as in the other score variants except that the fat ratio was replaced by olive oil use, cereals by whole grains, meat by red/processed meat, ethanol by red wine and fruits and nuts/seeds were separated into two components. Similar variants of the MDS have been used in other studies\(^{(27–30)}\).

Olive oil use was added as a separate component since it was not included in the nutrient calculations for fat, due to the design of the questionnaire in the CAPS study (yes/no, no frequency reported). The study-specific median intake in g/d was used as the cut-off between low and high intake of each component. The total MDS-alt ranging from zero to ten points was categorised into low (zero to three points), medium (four to five points) or high (six to ten points) adherence to the Mediterranean diet.

**Statistical methods**

We excluded men with incomplete dietary data (\(n\) 67) or unreasonably high or low energy intakes (<3300 kJ/d or >21 000 kJ/d) (\(n\) 27). In total, 2590 participants (1482 cases and 1108 controls) were included in the final analyses. Baseline characteristics of cases and controls were compared using the Wilcoxon–Mann–Whitney test for continuous variables and the \(\chi^2\) test for categorical variables. Correlations between the continuous MDS variants and their components were evaluated by estimating Spearman correlation coefficients.

Unconditional logistic regression was used to generate OR with 95 % CI, as estimates of the relative risk of prostate cancer according to adherence to the Mediterranean diet. Adherence to each MDS variant was analysed using indicator variables for low, medium and high score, with low as the reference group. Additionally, the scores were modelled as continuous variables, firstly by assessing the linear effect of a one-point increment in the score on prostate cancer risk and secondly by fitting restricted cubic splines to relax the linear assumption and to create smoothed functions of adherence to the scores. Moreover, we evaluated the individual effects of each MDS component. All analyses were performed for total prostate cancer, for advanced (\(n\) 588) and localised (\(n\) 512) disease, as well as for Gleason score 7 disease subtype (\(n\) 218).

Simple regression models included the matching factors: age and region of residence. Multivariate regression models additionally included education, smoking, BMI, energy intake, physical activity, history of diabetes and family history of prostate cancer. Potential confounders were selected based on subject matter knowledge as well as on indications provided by purely statistical procedures, such as a change in \(\beta\)-coefficients (>10 %). Other covariates that were considered as potential confounders but not included in the final model were height, employment status, marital status, snuff use, use of dietary supplements, use of olive oil and intake of coffee, phyto-oestrogens, potatoes, sweet foods and non-alcoholic beverages. The potential effect measure modification by selected covariates (age, family history of prostate cancer, history of diabetes and BMI) was evaluated in two ways: formal interaction tests by including multiplicative interaction terms in logistic regression models, and by the use of interaction indicator variables to obtain a stratified effect.

Wald and likelihood ratio tests were used to assess the statistical significance of observed associations (\(P<0.05\), two-sided tests). The fit of the models was evaluated by the Hosmer–Lemeshow and the Pearson \(\chi^2\) goodness-of-fit tests. All analyses were performed using the statistical software systems SAS version 9.2 (SAS Institute Inc.) and Stata version 12 (StataCorp LP).

**Results**

The study population is described in Table 2. The cases were somewhat younger than the controls and were more likely to reside in the northern part of Sweden, to be employed, to take dietary supplements and to have a family history of prostate cancer. Cases and controls also differed with regard to their intake of total energy, alcohol, non-alcoholic beverages, tomatoes and sweet foods, but there were no major differences concerning BMI, level of education, marital status, smoking status, physical activity, coffee intake or potato intake. The differences observed for the matching factors age and region of residence may be explained by the lower participation rate among controls than among cases, and was apparent already before exclusion of participants (\(n\) 94) (results not shown).

The mean total score of each of the five MDS variants, as well as the distribution between the three adherence groups, is given in Table 3. The results were in general similar for cases and controls, except for a slightly higher mean of MDS-serv, MDS-cent and MDS-alt among the cases. The mean total score of MDS-gram was 4.4 (SD 1.7) among both cases and controls: 31 % of cases and 33 % of controls had low adherence, 42 % of cases and 40 % of controls had medium adherence and 27 % of both cases and controls had high adherence.

Table 4 presents the intake levels of the MDS components and of the total energy, macronutrients and selected food items, in the CAPS population as well as in the Greek reference population. The intake of vegetables and fruits was considerably lower in the Swedish population, whereas the intake of legumes, cereals, fish and dairy products was higher than that in the Greek population. The intake of meat was similar in both populations. As expected, the MPS ratio in the
Table 2. Characteristics of the cases and controls in the Cancer of the Prostate in Sweden (CAPS) study

| Characteristics                        | Cases (n 1482) | Controls (n 1108) | P*   |
|----------------------------------------|---------------|------------------|------|
| **Age (years)**                        |               |                  |      |
| Mean                                   | 66.8          | 67.7             | <0.01|
| SD                                     | 7.3           | 7.5              |      |
| **Region of residence**                |               |                  | <0.01|
| Northern Sweden                        | 455           | 197              |      |
| Central Sweden                         | 1027          | 911              |      |
| **Education level**                    |               |                  | 0.15 |
| 0–9 years                              | 675           | 506              |      |
| 10–12 years                            | 595           | 468              |      |
| ≥13 years                              | 207           | 127              |      |
| **Marital status**                     |               |                  | 0.25 |
| Married/partner                        | 1200          | 867              |      |
| Divorced/unmarried                     | 204           | 165              |      |
| Widower                                | 77            | 71               |      |
| **History of diabetes**                |               |                  | 0.62 |
| Yes                                    | 171           | 134              |      |
| No                                     | 1292          | 953              |      |
| **Family history of prostate cancer**  |               |                  | <0.01|
| Yes                                    | 273           | 103              |      |
| No                                     | 1209          | 1005             |      |
| **Smoking**                            |               |                  | 0.44 |
| Never smoker                           | 575           | 424              |      |
| Former smoker                          | 734           | 535              |      |
| Current smoker                         | 156           | 134              |      |
| **Dietary supplement use**             |               |                  | <0.01|
| Yes                                    | 702           | 437              |      |
| No                                     | 737           | 623              |      |
| **BMI at inclusion (kg/m²)**           |               |                  | 0.44 |
| <25                                    | 574           | 400              |      |
| 25 to <30                              | 686           | 526              |      |
| ≥30                                    | 177           | 141              |      |
| **Physical activity (MET-h/d)**‡       |               |                  | 0.94 |
| Mean                                   | 12.7          | 12.8             |      |
| SD                                     | 8.9           | 9.1              |      |
| **Total energy intake (kJ/d)**         |               |                  | <0.01|
| Mean                                   | 9583.5        | 9303.9           |      |
| SD                                     | 2670.2        | 2721.1           |      |
| **Alcohol intake (g/d)**               |               |                  | <0.01|
| Mean                                   | 8.2           | 8.0              |      |
| SD                                     | 16.8          | 19.9             |      |
| **Non-alcoholic beverage intake (servings/d)** |           |                  | <0.01|
| Mean                                   | 1.0           | 0.9              |      |
| SD                                     | 1.2           | 1.2              |      |
| **Coffee intake (cups/d)**             |               |                  | 0.67 |
| Mean                                   | 3.1           | 3.1              |      |
| SD                                     | 1.9           | 2.0              |      |
| **Tomato intake (servings/d)**         |               |                  | <0.01|
| Mean                                   | 0.4           | 0.4              |      |
| SD                                     | 0.3           | 0.3              |      |
| **Potato intake (servings/d)**         |               |                  | 0.19 |
| Mean                                   | 0.5           | 0.5              |      |
| SD                                     | 0.3           | 0.3              |      |
| **Sweet foods intake (servings/d)**§   |               |                  | <0.01|
| Mean                                   | 0.8           | 0.8              |      |
| SD                                     | 0.5           | 0.5              |      |
| **Disease characteristics among cases (% of all cases)**| | | |
| Advanced]                              | 588           | 40               |      |
| Localised[‡]                           | 512           | 34               |      |
| Gleason score 7                        | 218           | 15               |      |
| Unknown                                | 87            | 6                |      |
| Other                                  | 77            | 5                |      |
| **Gleason score**                      |               |                  |      |
| Mean                                   | 6.5           |                  |      |
| SD                                     | 1.2           |                  |      |

Continued
Swedish population was lower than the MUFA:SFA ratio in the Greek population. The intake of SFA was similar in both populations, whereas the intake of MUFA and PUFA, as well as the total energy, was lower in the Swedish population. The Swedish population ate more potatoes and sweet foods but drank less non-alcoholic beverages than the Greek population. The correlation between individual components and the main score, MDS-gram, was low to moderate, ranging from \( r = 0.07 \) for ethanol to \( r = 0.58 \) for vegetables. Similarly, the individual components of the alternative score, MDS-alt, were weakly or moderately correlated with the total score (range \( r = 0.05-0.46 \)). Inter-correlation between the individual components was in the range \( r = 0.02-0.40 \) (results not shown). The correlation coefficients between the MDS variants ranged from 0.43 between MDS-serv and MDS-greek, to 0.79 between MDS-gram and MDS-serv and between MDS-gram and MDS-alt, respectively (results not shown).

As shown in Fig. 1, no statistically significant association was found between any of the MDS variants and prostate cancer. However, high MDS-cent, MDS-greek and MDS-alt scores seemed to be associated with a 20–35 % increased relative risk of advanced prostate cancer, compared with low scores, and a somewhat weaker increased risk of total prostate cancer. However, the estimates were not statistically significant and the CI were wide. For cases with Gleason score 7, a non-statistically significant inverse association was seen for high \( \nu \)s. Low MDS-gram and MDS-serv (multivariate OR were 0.84, 95 % CI 0.59, 1.19; and 0.77, 95 % CI 0.55, 1.09, respectively; results not shown). No association was seen between Gleason score 7 disease subtype and any of the other MDS-variants. There was no evidence of an association between the continuous scores and prostate cancer, in neither logistic regression models nor restricted cubic spline regression models, except for a non-significant positive association between the total prostate cancer and MDS-cent (results not shown).

Among the individual score components, a high vegetable intake was associated with an increased risk of prostate cancer, with multivariate OR 1.25 (95 % CI 1.05, 1.50; \( P_{\text{trend}} = 0.01 \)) for total prostate cancer, 1.37 (95 % CI 1.09, 1.72; \( P_{\text{trend}} = 0.006 \)) for advanced disease and 1.25 (95 % CI 0.98, 1.60; \( P_{\text{trend}} = 0.07 \)) for localised disease (results not shown). In supplementary analyses, the positive association remained but was weakened when tomatoes were excluded from vegetable intake, and a high intake of tomato products was found to be positively associated with localised disease (multivariate OR 1.25; 95 % CI 0.99, 1.59; \( P_{\text{trend}} = 0.07 \)) compared with low intake. A high ethanol intake was associated with an increased risk of advanced prostate cancer (multivariate OR 1.25; 95 % CI 0.99, 1.58; \( P_{\text{trend}} = 0.06 \)). Simple models

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### Table 2. Adherence to the Mediterranean Diet Score (MDS) variants in cases and controls in the Cancer of the Prostate in Sweden (CAPS) study* (Mean values and standard deviations; number of participants and percentages)

| Characteristics            | Cases (n 1482) | Controls (n 1108) |
|----------------------------|---------------|------------------|
|                           | n  | %†  | n  | %†  | P*  |
| **PSA level (ng/ml)**      |    |     |    |     |     |
| Mean                       | 48 | 33  | 46 | 33  |     |
| SD                         | 82 | 57  | 75 | 53  |     |
| **MDS-gram (points)**      |    |     |    |     |     |
| Mean                       | 4  | 27  | 1  | 9   |     |
| SD                         | 1  | 8   | 1  | 8   |     |
| **MDS-serv (points)**      |    |     |    |     |     |
| Mean                       | 4  | 27  | 1  | 9   |     |
| SD                         | 1  | 8   | 1  | 8   |     |
| **MDS-cent (points)**      |    |     |    |     |     |
| Mean                       | 4  | 27  | 1  | 9   |     |
| SD                         | 1  | 8   | 1  | 8   |     |
| **MDS-greek (points)**     |    |     |    |     |     |
| Mean                       | 4  | 27  | 1  | 9   |     |
| SD                         | 1  | 8   | 1  | 8   |     |
| **MDS-alt (points)**       |    |     |    |     |     |
| Mean                       | 4  | 27  | 1  | 9   |     |
| SD                         | 1  | 8   | 1  | 8   |     |

* For details of the MDS variants, see Table 1.
† Percentage of all cases and of all controls, respectively.

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### Table 3. Adherence to the Mediterranean Diet Score (MDS) variants in cases and controls in the Cancer of the Prostate in Sweden (CAPS) study* (Mean values and standard deviations; number of participants and percentages)

| Characteristics            | Cases (n 1482) | Controls (n 1108) |
|----------------------------|---------------|------------------|
|                           | n  | %†  | n  | %†  | P*  |
| **MDS-gram (points)**      |    |     |    |     |     |
| Mean                       | 4  | 27  | 1  | 9   |     |
| SD                         | 1  | 8   | 1  | 8   |     |
| **MDS-serv (points)**      |    |     |    |     |     |
| Mean                       | 4  | 27  | 1  | 9   |     |
| SD                         | 1  | 8   | 1  | 8   |     |
| **MDS-cent (points)**      |    |     |    |     |     |
| Mean                       | 4  | 27  | 1  | 9   |     |
| SD                         | 1  | 8   | 1  | 8   |     |
| **MDS-greek (points)**     |    |     |    |     |     |
| Mean                       | 4  | 27  | 1  | 9   |     |
| SD                         | 1  | 8   | 1  | 8   |     |
| **MDS-alt (points)**       |    |     |    |     |     |
| Mean                       | 4  | 27  | 1  | 9   |     |
| SD                         | 1  | 8   | 1  | 8   |     |

* For details of the MDS variants, see Table 1.
† Percentage of all cases and of all controls, respectively.
Table 4. Summary statistics of the intake of the components of the Mediterranean Diet Score (MDS) variants, of the intake of energy, macronutrients and selected food items, and their correlation with the total score, for the controls of the Cancer of the Prostate in Sweden (CAPS) study and for the male study population in the European Prospective Investigation into Cancer and Nutrition (EPIC) in Greece (Twenty-fifth centiles; median values; mean values and standard deviations; seventy-fifth centiles; Spearman correlation coefficients)

| Components in MDS-gram, MDS-serv, MDS-cent and MDS-greek | Twenty-fifth centile | Median | Mean | SD | Seventy-fifth centile | Spearman r‡ |
|----------------------------------------------------------|----------------------|--------|------|----|----------------------|------------|
| Ratio of MUFA and PUFA to SFA                            | 1.0                  | 1.3    | 1.4  | 0.5 | 1.7                  | 0.49       |
| Vegetables (g/d)                                         | 80.3                 | 118.2  | 137.8| 83.5| 176.6                | 0.58       |
| Fruits, nuts and seeds (g/d)                            | 70.1                 | 116.0  | 131.7| 83.8| 176.7                | 0.50       |
| Legumes (g/d)                                            | 15.8                 | 26.2   | 32.8 | 27.5| 41.4                 | 0.46       |
| Cereals (g/d)                                            | 219.0                | 278.8  | 304.5| 125.7| 377.2                | 0.35       |
| Fish (g/d)                                               | 25.6                 | 36.5   | 41.5 | 24.2| 51.5                 | 0.32       |
| Dairy products (g/d)                                     | 344.8                | 551.8  | 612.1| 366.1| 820.4                | -0.34      |
| Meat/meat products (g/d)                                 | 90.2                 | 120.1  | 127.0| 52.8| 156.3                | -0.09      |
| Ethanol (g/d)                                            | 1.4                  | 4.1    | 8.2  | 15.1| 8.6                  | 0.07       |

**Additional components in MDS-alt**

| Olive oil use (yes/no)                                   | -                    | -      | -    | -   | -                    | -0.48      |
| Fruits (g/d)                                             | 68.8                 | 115.2  | 130.4| 83.3| 175.0                | 0.46       |
| Nuts and seeds (g/d)                                     | 0                    | 0.1    | 1.3  | 2.4 | 1.8                  | 0.46       |
| Whole grains (g/d)                                       | 75.4                 | 121.7  | 145.4| 100.5| 190.0                | 0.26       |
| Red/processed meat (g/d)                                 | 78.5                 | 107.5  | 113.5| 49.6| 141.4                | -0.05      |
| Red wine (g/d)                                           | 0                    | 0      | 28.9 | 46.3| 41.8                 | -0.33      |

**Intake of energy, macronutrients, and selected food items not included in the MDS variants**

| Total energy (kJ/d)                                      | 7300.1               | 8930.7 | 9303.9| 2721.1| 10925.6               | -0.33      |
| Total fat (g/d)                                          | 60.7                 | 76.0   | 81.3 | 29.5 | 97.4                  | -0.35      |
| Saturated fat (g/d)                                      | 25.7                 | 33.2   | 36.0 | 14.4 | 43.9                  | -0.43      |
| Monounsaturated fat (g/d)                                | 21.0                 | 26.2   | 28.2 | 10.2 | 33.3                  | -0.31      |
| Polyunsaturated fat (g/d)                                | 8.0                  | 9.9    | 10.7 | 4.2  | 12.7                  | -0.08      |
| Carbohydrates (g/d)                                      | 214.8                | 263.4  | 270.8| 81.7 | 315.4                 | -0.24      |
| Protein (g/d)                                            | 68.6                 | 83.5   | 86.0 | 24.5 | 98.9                  | -0.25      |
| Potatoes (g/d)                                           | 62.5                 | 108.2  | 106.3| 62.1 | 146.9                 | 0.03       |
| Sweet foods (g/d)**                                      | 21.8                 | 37.8   | 43.5 | 29.7 | 59.4                  | 0.05       |
| Non-alcoholic beverages (g/d)**                          | 0                    | 133.9  | 207.4| 263.0| 294.8                 | -0.07      |

| Components of the original MDS†                          |                      |        |      |     |                      |            |
| Ratio of MUFA to SFA                                     | 1.7                  | 1.8    | 0.5  |     |                      |            |
| Vegetables (g/d)                                         | 549.9                | 583.6  | 233.9|     |                      |            |
| Fruits and nuts (g/d)                                    | 362.5                | 393.0  | 214.6|     |                      |            |
| Legumes (g/d)                                            | 9.1                  | 10.4   | 7.4  |     |                      |            |
| Cereals (g/d)                                            | 266.4                | 191.0  | 80.2 |     |                      |            |
| Fish (g/d)                                               | 23.7                 | 26.4   | 20.3 |     |                      |            |
| Dairy products (g/d)                                     | 196.7                | 222.6  | 147.6|     |                      |            |
| Meat/meat products (g/d)                                 | 120.8                | 129.3  | 60.4 |     |                      |            |
| Ethanol§                                                 | -                    | -      | -    |     |                      |            |

| Intake of components not included in the original MDS    |                      |        |      |     |                      |            |
| Total energy (kJ/d)                                      | 9851.4               | 10202.6| 2949.3|     |                      |            |
| Saturated fat (g/d)                                      | 33.1                 | 34.6   | 13.2 |     |                      |            |
| Monounsaturated fat (g/d)                                | 55.9                 | 58.4   | 20.0 |     |                      |            |
| Polyunsaturated fat (g/d)                                | 15.0                 | 17.5   | 9.2  |     |                      |            |
| Carbohydrates (g/d)                                      | 314.5                | 395.4  | 271.6|     |                      |            |
| Protein (g/d)                                            | 68.9                 | 83.5   | 86.0 | 24.5 | 98.9                  | -0.25      |
| Potatoes (g/d)                                           | 88.7                 | 98.9   | 63.3 |     |                      |            |
| Sweets (g/d)**                                           | 22.8                 | 26.8   | 19.7 |     |                      |            |

* All intakes in the CAPS study except energy intake are adjusted to a daily total energy intake of 10 460 kJ (equivalent to 2500 kcal). For details of the MDS variants, see Table 1.

† Trichopoulou et al(20).
‡ Correlation with the total score (MDS-gram), except where marked §, whose definition is given later.
§ Categorisation of ethanol (<10 g/d; 10 to <30 g/d; ≥30 g/d) was independent of the median intake in the study population.
∥ Correlation with total score (MDS-alt).
¶ Not included as a component of the MDS in the EPIC study.
** Includes confectionery, sweet bakery products and ice cream.
†† Includes juices and soft drinks.
showed an inverse association for a high MP:S ratio, but the association disappeared after adjusting for potential confounders. A high meat intake was associated with a 10–15 % reduction in prostate cancer risk in multivariable models, although this was not statistically significant. No association was found between the intake of fruits and nuts, legumes, cereals, fish or dairy products and prostate cancer. The results for Gleason score 7 disease subtype were in the same direction as for total prostate cancer (results not shown). As illustrated in Fig. 2, smoothed functions of intake showed weak positive associations between total prostate cancer and the MP:S ratio, vegetable intake and fruit/nut intake, although the CI included the value of 1. A dose–response trend was suggested for the association with the MP:S ratio. A high intake of legumes, fish and meat was possibly associated with a non-significant reduced risk of prostate cancer, whereas no association was observed for cereals, dairy products or alcohol.

Analysis of components specific to the MDS-alt showed an inverse association between a high intake of red/processed meat and total prostate cancer (multivariable OR 0.83; 95 % CI 0.70, 0.99; $P$ trend = 0.04). The results were similar but non-statistically significant in the case of advanced and localised disease. We observed an increased risk of advanced disease for a high intake of red wine (multivariable OR 1.39; 95 % CI 1.06, 1.82; $P$ trend = 0.02). No statistically significant associations were seen for olive oil use, whole grains, fruit or nuts.

Tests for potential interactions between the Mediterranean diet (MDS-gram) and selected covariates showed no statistically significant interactions (results not shown). However, when data were stratified using interaction indicator variables, men with a family history of prostate cancer had a non-significant increased risk of total prostate cancer (OR 1.67; 95 % CI 0.87, 3.22; $P_{interaction}$ = 0.29) comparing high with low MDS-gram, which was not seen in men without a family history of the disease; this possible interaction was even more pronounced for advanced disease (OR 2.42; 95 % CI 1.10, 5.34; $P_{interaction}$ = 0.10). Furthermore, stratified analyses showed a positive association between the Mediterranean diet and prostate cancer among obese men (BMI $\geq$ 30 kg/m²), with multivariable OR 2.06 (95 % CI 1.11, 3.81; $P_{interaction}$ = 0.09) for total malignancy, 2.17 (95 % CI 0.99, 4.76; $P_{interaction}$ = 0.24) for advanced disease and 1.94 (95 % CI 0.84, 4.49; $P_{interaction}$ = 0.48) for localised disease, but not among men with BMI $<$ 30 kg/m². Age at inclusion and diabetes did not act as effect modifiers in the present study.

**Discussion**

Overall, we found no association between the Mediterranean diet and relative risk of prostate cancer. Among individual score components, a high intake of vegetables and alcohol was associated with an increased risk of prostate cancer, and a high intake of red/processed meat with a decreased risk of

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### Table: Adherence to the Mediterranean Diet Score (MDS) and Prostate Cancer Risk

| Component          | Adherence Score | OR   | 95 % CI |
|--------------------|-----------------|------|---------|
| Total PC           | Medium          | 1.08 | 0.68, 1.33 |
|                   | High            | 1.03 | 0.61, 1.30 |
| Advanced PC        | Medium          | 1.09 | 0.91, 1.13 |
|                   | High            | 1.09 | 0.81, 1.38 |
| Localised PC       | Medium          | 1.50 | 1.05, 2.13 |
|                   | High            | 1.08 | 0.76, 1.50 |

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### Fig. 1. Relative risk of total (n = 2336), advanced (n = 1494) and localised (n = 1441) prostate cancer (PC) according to adherence to the Mediterranean Diet Score (MDS) variants: (a) MDS-gram, (b) MDS-serv, (c) MDS-cent, (d) MDS-greek and (e) MDS-alt. For details of the MDS variants, see Table 1. Estimates reflect OR (95 % CI) derived from unconditional logistic regression models. All models are adjusted for age (in 5-year intervals), region of residence (north/central), education (0–15 years; ≥ 16 years), smoking status (never/former/current), BMI (quartile distribution of controls), energy intake (quartile distribution of controls), physical activity (quartile distribution of controls), history of diabetes (yes/no) and family history of prostate cancer (yes/no). Adherence groups were categorised as follows: (a–c) low adherence, 0–3 points (reference); medium adherence, 4–5 points; high adherence, 6–9 points; (d) low adherence, 0–2 points (reference); medium adherence, 3–4 points; high adherence, 5–9 points; (e) low adherence, 0–3 points (reference); medium adherence, 4–5 points; high adherence, 6–10 points.
the disease. The association between MDS and prostate cancer was indicated as positive in obese men and in men with a family history of the disease.

To the best of our knowledge, no other studies have so far been carried out on the association between the Mediterranean dietary pattern and prostate cancer using a dietary score. A few studies looking at empirically derived dietary patterns have shown an increased risk of prostate cancer with ‘Western’ dietary patterns rich in meat and processed foods(31–33), i.e. foods consumed at low amounts in the traditional Mediterranean diet. Interestingly, Greek migrants in Australia have a lower risk of prostate cancer than men born in Australia, and they have also retained the dietary habits of their native country, as opposed to many other migrant populations(4). Studies on the Mediterranean diet and overall cancer risk have shown inconsistent results(2,3). A meta-analysis of data from seven cohort studies provided a pooled relative risk of cancer of 0.94 (95% CI 0.92, 0.96) for a two-point increase in the MDS(5). In a recent review, the Mediterranean diet was considered as ‘probably’ protective against overall cancer, albeit there is variation with regard to different cancer sites(3).

Although the Mediterranean diet as a whole has not previously been studied in relation to prostate cancer, its individual components have. Recent reviews suggest that a high intake of vegetables, marine fatty acids, legumes and foods containing vitamin E (such as vegetable oils, nuts and seeds) may offer protection against the disease(4,34–37), and a favourable effect may also be seen with a low intake of meat and foods rich in Ca such as dairy products(4,35–38). In theory, the combined effect of these components may be beneficial in prostate cancer prevention.

In the present study, the observed positive association between a high vegetable intake and prostate cancer is unexpected, considering earlier indications of an inverse association(34,36). The main contributors to the vegetable intake were tomatoes, carrots and mixed vegetables, each contributing on average 14–16%. A high intake of tomatoes, notably tomato sauce, has been shown in several studies to reduce the risk of prostate cancer(34,37,39). The indicated positive association between tomato intake and localised disease in our study may be explained by localised cases potentially being more health conscious and more likely to eat healthier.

Fig. 2. Dose–response relationship between components of the Mediterranean Diet Score (MDS-gram; see Table 1 for details) and relative risk of total prostate cancer (n = 2336): (a) MUFA and PUFA to SFA (MP:S) ratio, (b) vegetables, (c) fruits and nuts, (d) legumes, (e) cereals, (f) fish, (g) meat and meat products, (h) dairy products and (i) alcohol. Results from restricted cubic regression splines with five knots. (—), OR; (---), 95% CI. The CI cross at the reference point, equivalent to the tenth centile of the intake in the study population. Regression models include age (in 5-year intervals), region of residence (north/central), education (0–9 years; 10–12 years; ≥13 years), smoking status (never/former/current), BMI (quartile distribution of controls), energy intake (quartile distribution of controls), physical activity (quartile distribution of controls), history of diabetes (yes/no) and family history of prostate cancer (yes/no).
The increased risk for the total vegetable intake was strongest for advanced disease and remained, although weakened, when tomatoes were excluded from the vegetable intake. Also, the vegetable intake was much lower in the study population than in the Greek reference population, and thus a true detrimental effect is unlikely. We also observed positive associations between advanced disease and intakes of total alcohol and red wine, although there is limited evidence for an association between alcohol and prostate cancer\(^{(36,37)}\). Moreover, our findings of an inverse association with red/processed meat and no association with dairy products are in contrast with previous findings of an increased risk of prostate cancer for high intakes of meat and Ca, although the evidence is inconsistent\(^{(36,37,41,42)}\). The deviating results in our study may be explained by residual confounding or recall bias, or may be due to chance, considering that we tested multiple hypotheses. Another component of the MDS that deserves attention is the fish intake. Previous studies on the intake of fatty fish rich in n-3 fatty acids have shown inverse associations with prostate cancer\(^{(43,44)}\), whereas a recent meta-analysis of the total fish intake showed no association\(^{(45)}\). The MDS does not distinguish between fatty and non-fatty fish, which may explain the lack of association in our study.

Olive oil is the main source of fat in the Mediterranean region and was included as a component of the alternative score, MDS-alt. It has been shown to protect against several types of cancer\(^{(46,47)}\), but no consistent association has been found with prostate cancer\(^{(48)}\). We observed no association between olive oil use and prostate cancer. Due to the generally low intake of olive oil in the Swedish population at the time of data collection, this component probably has little overall influence on the present results.

The construction of a dietary score as a measure of a healthy diet is complex, and the MDS is no exception. An inappropriate scoring method may be unable to detect a true association or may lead to misclassification of exposure. For instance, the score may be too blunt to detect a weak association with cancer since it does not capture all dietary aspects or factors unrelated to diet, or because the beneficial and non-beneficial components of the score may counter-balance each other’s potential effects, leading to a null result. This is a possible explanation for the overall lack of association seen in the present study. Also, strong correlations between the score and its components may influence the discriminatory power of each individual component. In the present study, the estimated inter-correlation between all components and their correlation with the total score was low to moderate, which means that each component contributes equally to the score. An exception was the vegetable component with a relatively high correlation with the total score (r=0.58); hence, it may be the dominant factor determining the score. Méa-Villarroya et al.\(^{(49)}\) recently evaluated the reliability of several indexes assessing adherence to the Mediterranean diet, and the MDS showed high performance.

Using study-specific intakes, as opposed to externally defined values, as cut-off between low and high intake increases the statistical power in each intake group, but it does not necessarily mean that the score is able to discriminate between beneficial and non-beneficial health effects. This may be problematic especially in non-Mediterranean populations where intake levels may be very different from those of a typical Mediterranean population. The main differences between our Swedish study population and the Greek reference population were the considerably lower intake of fruit and vegetables and higher dairy product intake in the former. However, the two populations differ not only in nationality but also in age, the CAPS population being older, and in the type of exposure data, the Greek data being longitudinal and prospective whereas the Swedish data were collected only once and retrospectively. Nevertheless, the observed differences in intake could indicate a limited discriminatory power of the MDS in the study population of the present study.

To explore the influence of using study-specific intakes as cut-off values, we compared variants of the MDS with different cut-off values, including one score based on the median intake in a Greek reference population. Overall, we did not find any statistically significant association with prostate cancer for any of the score variants. The non-significant increased risk of disease found with MDS-cent and MDS-greek was unexpected, as we had rather anticipated inverse associations. These results are probably driven by a high vegetable intake, as the vegetable component had the highest correlation with the total score and was positively associated with prostate cancer risk, especially advanced disease. Since overall we observed no major differences between the score variants, we suggest that the use of study-specific intakes as cut-off values in the MDS is appropriate also in non-Mediterranean populations. The usefulness of the MDS in non-Mediterranean populations is further supported by an Australian study where the MDS was found to be appropriate for the prediction of survival in both a population of Mediterranean descent (Greek-Australians) and a non-Mediterranean population (Anglo-Celts)\(^{(50)}\). Also, in a European multi-centre cohort study, a two-point increment in the MDS yielded risk estimates for overall cancer that were similar in men in Northern and Southern European countries\(^{(51)}\).

We also created an alternative score variant, MDS-alt, to investigate the effect of a more specific score that better reflects the ‘original’ Mediterranean diet. Replacing the fat ratio with olive oil use, cereals with whole grains, meat with red/processed meat and ethanol with red wine, as well as separating fruits from nuts, ideally captures the most health-beneficial components of the Mediterranean diet. Comparing MDS-gram and MDS-alt, which both used study-specific median intakes as cut-off values, we found no statistically significant associations for either of them, the two scores being highly correlated. The non-statistically significantly increased relative risk of advanced prostate cancer seen with MDS-alt is surprising, and may be driven by the vegetable component as previously discussed, or may be due to residual confounding.

The influence of diet on risk of developing prostate cancer may differ between men who are diagnosed in young or old age, or between men with and without a family history of the disease, since the type of prostate cancer and disease mechanisms may differ between the groups. We also hypothesised that diabetic men, as well as overweight or obese men, may have different metabolic responses to a Mediterranean diet compared with non-diabetic or normal-weight men. Formal
interaction tests showed no statistically significant interactions for any of these factors. However, in stratified analyses we observed positive associations between adherence to the Mediterranean diet and prostate cancer among men with a family history of prostate cancer and among obese men, but not among men without a family history of the disease or who were non-obese. These results should be interpreted with caution, especially since the control groups were relatively small (89 for those with a family history of prostate cancer and 129 for BMI ≥ 30 kg/m²).

The strengths of our study include the large sample size, high response-rate, the use of population-based random controls, complete and rapid ascertainment of prostate cancer cases and clinical information on prostate cancer subtypes. The proportion of prostate-specific antigen-detected cases was low (29% of 1499) due to the low frequency of prostate-specific antigen testing at the time of enrolment (14), ensuring a high proportion of cases with clinically relevant prostate cancer. The risk of confounding by population stratification was minimised by the ethnic homogeneity of the study population. Also, the extensive questionnaire provided detailed information on both exposure and potential confounding factors. Selection bias is possible since cases are generally more prone to participate than controls, and selection of health-conscious controls may occur. The retrospective collection of exposure information may produce recall bias. However, a previous study comparing original and repeated dietary recall interviews showed very little or no overall difference in performance between prostate cancer cases and non-cases (52). Another drawback is the fact that the FFQ assesses dietary intake 1 year prior to diagnosis, which may differ from dietary habits earlier in life, before tumour initiation. Furthermore, reverse causation bias cannot be ruled out, although men habits earlier in life, before tumour initiation. Furthermore, reverse causation bias cannot be ruled out, especially since the control groups were relatively small (89 for those with a family history of prostate cancer and 129 for BMI ≥ 30 kg/m²).

The potential effect of any remaining measurement error on our relative risk estimates is attenuation, which would reduce the likelihood of detecting a potentially weak diet–disease relationship (57); thus it may in part explain the lack of an association seen between the Mediterranean diet and prostate cancer.

In conclusion, we found no support for any association between the Mediterranean dietary pattern and prostate cancer in the Swedish CAPS study. We have compared several MDS variants with different compositions and cut-off values, and observed no significant associations with any of the scores. We suggest that using study-specific median intakes as cut-off values as in the original MDS is an appropriate way of assessing adherence to the Mediterranean diet also in non-Mediterranean populations. However, the present results, including the unexpected associations seen for certain components of the MDS, need to be cautiously interpreted in light of potential limitations in the case–control design, the exposure assessment or the discriminatory power of the MDS in our study population. Further studies are needed to clarify the relationship between the Mediterranean diet and prostate cancer risk.

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References

1. Ferlay J, Shin HR, Bray F, et al. (2010) GLOBOCAN 2008 Cancer Incidence and Mortality Worldwide: Version 1.2. IARC CancerBase
48. Psaltopoulou T, Kosti RI, Haidopoulos D, et al. (2011) Olive oil intake is inversely related to cancer prevalence: a systematic review and a meta-analysis of 13 800 patients and 23 340 controls in 19 observational studies. *Lipids Health Dis* 10, 127.

49. Mila-Villarroel R, Bach-Faig A, Puig J, et al. (2011) Comparison and evaluation of the reliability of indexes of adherence to the Mediterranean diet. *Public Health Nutr* 14, 2338–2345.

50. Kouris-Blazos A, Gnardellis C, Wahlqvist ML, et al. (1999) Are the advantages of the Mediterranean diet transferable to other populations? A cohort study in Melbourne, Australia. *Br J Nutr* 82, 57–61.

51. Couto E, Boffetta P, Lagiou P, et al. (2011) Mediterranean dietary pattern and cancer risk in the EPIC cohort. *Br J Cancer* 104, 1493–1499.

52. Wilkens LR, Hankin JH, Yoshizawa CN, et al. (1992) Comparison of long-term dietary recall between cancer cases and noncases. *Am J Epidemiol* 136, 825–835.

53. Demark-Wahnefried W, Peterson B, McBride C, et al. (2000) Current health behaviors and readiness to pursue life-style changes among men and women diagnosed with early stage prostate and breast carcinomas. *Cancer* 88, 674–684.

54. Mroz IW, Chapman GE, Oliffe JL, et al. (2010) Men, food, and prostate cancer: gender influences on men’s diets. *Am J Mens Health* 5, 177–187.

55. Patterson RE, Neuhouser ML, Heddersen MM, et al. (2003) Changes in diet, physical activity, and supplement use among adults diagnosed with cancer. *J Am Diet Assoc* 103, 323–328.

56. Nordic Council of Ministers (2004) *Nordic Nutrition Recommendations 2004: Integrating Nutrition and Physical Activity*, 4th ed. Copenhagen, Denmark: Nordic Council of Ministers.

57. Willett W (1998) *Nutritional Epidemiology*, 2nd ed. New York: Oxford University Press.

58. Willett WC (1994) Future directions in the development of food-frequency questionnaires. *Am J Clin Nutr* 59, Suppl. 1, S171–S174.

59. Benitez-Arciniega AA, Mendez MA, Baena-Diez JM, et al. (2011) Concurrent and construct validity of Mediterranean diet scores as assessed by an FFQ. *Public Health Nutr* 14, 2015–2021.