OBJECTIVE  To examine stage-matched nutrition counseling by family physicians and its effect on dietary intake, anthropometry, and serum lipid levels in patients at elevated risk for cardiovascular disease.

METHODS In this controlled trial, patients randomized to intervention practices received nutrition information following the Stages-of-Change Model, and patients randomized to control practices received usual care.

RESULTS At both 6 and 12 months after baseline, total fat intake and saturated fat intake declined significantly more in the intervention group than in the control group: -5.7% and -2.6% of energy, respectively, at 6 months, and -3.6% and -1.7% of energy, respectively, at 12 months. For energy intake, body weight, and BMI, there were significant differences between groups only at 6 months: -0.8 megajoules (MJ), -0.7 kg, and -0.3 kg/m², respectively. None of the serum lipid values changed significantly between groups at 12 months.

CONCLUSIONS Nutritional counseling based on stages of change led to reductions in dietary fat intake and weight loss in the short term. However, we found no corresponding changes in serum lipid concentrations.

KEY WORDS  Nutrition counseling; models of change; serum lipids; cardiovascular disease. (J Fam Pract 2002; 51:751–758)
However, in these studies the intervention was not managed by the FP. We think the FP is the most appropriate person to manage such intervention in the family practice. Therefore, we conducted a controlled dietary intervention, based on the Stages-of-Change Model and managed by the FP with selective referral to a dietician. We examined the effects of dietary counseling on changes in dietary intake, anthropometry, and serum lipid levels in patients at elevated cardiovascular risk.

METHODS

Participants and design

In this randomized controlled trial, men and women at elevated risk for cardiovascular disease were recruited from the 9 family practices joining the Nijmegen Monitoring Project, the research network of the Department of Family Medicine, University Medical Centre St. Radboud. Selection and flow of participants are described in Figure 1. Seventy-one patients were initially included in the intervention group and 72 patients in the control group (Table 1). Consequently, we had enough power (0.90) to detect a difference in change between groups for total fat intake of 3% of energy and 11.6 mg% of serum total cholesterol.

After selection and recruitment of participants, the family practices were randomly divided into intervention (4) and control (5) practices. The practices, and not the patients, were the units of randomization, to avoid contamination of the information between intervention and control groups. Patients in the control practices received usual care. Each patient in the intervention arm received nutrition information according to his or her stage of change. All participants signed an informed consent form before entering the study. The Medical Ethical Committee of the Department of Human Nutrition and Epidemiology, Wageningen University, approved the study protocol. The study lasted from August 1998 until April 2000.

Measurements

Specially trained practice assistants measured anthropometry data, and presented patients with a self-administered questionnaire on demographics, medical history, food frequency, and a stages-of-change algorithm at baseline, 6 months, and 12 months. Blood samples were taken at baseline and at 12 months.

Physical assessment. Anthropometry consisted of body weight to the nearest 0.5 kg, height, and waist and hip circumferences to the nearest 0.5 cm. Patients wore no shoes and only light clothing when weighed. Fasting blood samples were taken twice per measurement period at a 1-week interval, with the patient in the sitting position. The samples were stored at -80°C. Lipids were analyzed enzymatically for total cholesterol, HDL cholesterol, LDL cholesterol, and triglycerides with the Cobas Intergra 700 (Roche Diagnostics, Switzerland), at the laboratory of the Canisius Wilhelmina Hospital (Nijmegen, The Netherlands). The coefficient of variation within runs was 2.3% for total cholesterol, 1.6% for HDL cholesterol, and 1.8% for triglycerides. The LDL cholesterol level was calculated using the equation of Friedewald et al.

Questionnaires. The questionnaire asked for demographic data, family history of heart disease, smoking status, physical activity, drug use, and diet

<table>
<thead>
<tr>
<th>TABLE 1</th>
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<tbody>
<tr>
<td><strong>Baseline characteristics of patients in the intervention and control groups</strong></td>
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<tr>
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<tr>
<td>Sex (%)</td>
</tr>
<tr>
<td>Male</td>
</tr>
<tr>
<td>Female</td>
</tr>
<tr>
<td>Age (years)</td>
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<tr>
<td>Disorder (%)</td>
</tr>
<tr>
<td>Hypertension</td>
</tr>
<tr>
<td>Diabetes mellitus II</td>
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<tr>
<td>Hypertension &amp; diabetes mellitus II</td>
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<tr>
<td>Marital status (%)</td>
</tr>
<tr>
<td>Single</td>
</tr>
<tr>
<td>Married/cohabiting</td>
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<tr>
<td>Divorced</td>
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<tr>
<td>Widowed</td>
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<tr>
<td>Education (%)</td>
</tr>
<tr>
<td>Low</td>
</tr>
<tr>
<td>Intermediate</td>
</tr>
<tr>
<td>High</td>
</tr>
<tr>
<td>Family history of heart disease (%)</td>
</tr>
<tr>
<td>First-degree relatives younger than 60 years</td>
</tr>
<tr>
<td>Smoking (%)</td>
</tr>
<tr>
<td>Not smoking</td>
</tr>
<tr>
<td>Light smoker</td>
</tr>
<tr>
<td>Heavy smoker</td>
</tr>
<tr>
<td>Exercise, more than 20 minutes</td>
</tr>
<tr>
<td>No exercise</td>
</tr>
<tr>
<td>Less than 3 times a week</td>
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<tr>
<td>3 times a week</td>
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<tr>
<td>More than 3 times a week</td>
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</tbody>
</table>

1 Two-sided P values for differences in baseline characteristics between intervention and control groups.
3 First-degree relatives younger than 60 years.
4 Light smoker: 0-10 cigarettes a day, or smoking pipe or cigars. Heavy smoker: > 10 cigarettes a day.
history at baseline. At follow-up we checked for changes in smoking status, physical activity, and drug use. Patients in the control group were asked if they had visited a dietician during the study period.

The intake of energy, total fat, fatty acids, and cholesterol during the preceding 4 weeks was assessed by asking patients to fill out a food frequency questionnaire that included 104 food items. The questionnaire was validated and recently revised according to the Dutch National Food Survey. Dieticians carried out nutrient calculations with a computerized version of the Dutch food composition tables2 and

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**Selection and flow of participants**

**Study population:**
Patients from 9 general practices of the Nijmegen Monitoring Project*

n=46,500

Inclusion criteria:
- Type 2 diabetes mellitus or hypertension
- Age 40–70 years
- No manifest cardiovascular disease
- Not under treatment of a cardiologist
- Last measured serum total cholesterol ≥ 239.4 mg/dL, or unknown

n=416

Further selection in family practice:
- Dietary intake of total fat ≥ 37% or saturated fat ≥ 12%, expressed as percent of energy intake; and
- Mean serum total cholesterol ≥ 239.4 mg/dL based on 2 measurements with 1-week interval

n=400

Randomization of practices

**Intervention**
4 practices (n=71)

Nutrition counseling based on stages of change

n=70
Reason for drop out: don’t want to participate anymore (n=1)

n=69
Reasons for drop out: illness (n=1), don’t want to participate anymore (n=4)

**Control**
5 practices (n=72)

Usual care

n=67
Reasons for drop out: referral to cardiologist (n=1), don’t want to participate anymore (n=4)

n=63
Reason for drop out: don’t want to participate anymore (n=4)
phoned patients in cases of inconsistency.

Stages of change for reduction of fat intake were assessed with a 4-item algorithm based on measures used in previous studies in combination with the results of the food frequency questionnaire. According to the algorithm, participants were judged to be in precontemplation if they did not consider their diet to be low in fat, they were not in the process of cutting down on fat, and they had no intention of reducing their fat consumption. Participants were considered to be in the contemplation phase if they intended to decrease their fat intake within 6 months but not within 30 days, and to be in preparation when they intended to decrease their fat intake within 30 days. Participants who reported they were currently trying to eat less fat were classified as in action, and participants who reported they had been eating less fat for at least 6 months were classified in maintenance. If participants in maintenance consumed ≥ 37% total fat or ≥ 12% saturated fat expressed as percent of energy intake, they were reclassified in precontemplation.

After completing the study, all patients filled in an evaluation questionnaire. Patients in the control group were also asked about which nutrition information they had received during the last year.

**Intervention**

The intervention consisted of nutrition counseling based on stages of change, directed by the FP with selective referral to a dietician. FPs were supported by a protocol that included Prochaska’s stages of change.

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**TABLE 2**

Baseline measures and changes after 6 months and 12 months in dietary intake and anthropometry

<table>
<thead>
<tr>
<th></th>
<th>At Baseline</th>
<th></th>
<th>At 6 months</th>
<th></th>
<th>At 12 months</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Intervention</td>
<td>Control</td>
<td>P*</td>
<td>Intervention</td>
<td>Control</td>
<td>P**</td>
</tr>
<tr>
<td>n=71</td>
<td>Mean ± SD</td>
<td>n=72</td>
<td>Mean ± SD</td>
<td>n=70</td>
<td>Mean ± SD</td>
<td>n=67</td>
</tr>
<tr>
<td><strong>Dietary intake</strong> (per day)</td>
<td></td>
<td></td>
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<tr>
<td>Total energy (MJ/d)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>9.1 ± 2.7</td>
<td>9.6 ± 2.6</td>
<td>0.25</td>
<td>-1.4 ± 1.9t</td>
<td>-0.6 ± 1.8t</td>
<td>0.01</td>
<td>-0.7 ± 3.0t</td>
</tr>
<tr>
<td>Total fat (% of energy)</td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>42.1 ± 6.3</td>
<td>42.6 ± 5.2</td>
<td>0.64</td>
<td>-7.9 ± 6.5t</td>
<td>-2.2 ± 4.9t</td>
<td>0.00</td>
<td>-5.6 ± 6.9t</td>
</tr>
<tr>
<td>Saturated fat (% of energy)</td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>15.2 ± 2.6</td>
<td>15.5 ± 2.3</td>
<td>0.42</td>
<td>-3.4 ± 2.7t</td>
<td>-0.8 ± 2.2t</td>
<td>0.00</td>
<td>-2.6 ± 2.7t</td>
</tr>
<tr>
<td>Monounsaturated fat (% of energy)</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>14.6 ± 3.3</td>
<td>14.9 ± 2.6</td>
<td>0.53</td>
<td>-3.4 ± 3.3t</td>
<td>-0.7 ± 2.4t</td>
<td>0.00</td>
<td>-1.9 ± 4.1t</td>
</tr>
<tr>
<td>Unsaturated fat (% of energy)</td>
<td></td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>9.4 ± 3.0</td>
<td>9.3 ± 3.0</td>
<td>0.79</td>
<td>-1.0 ± 3.1t</td>
<td>-0.8 ± 3.0t</td>
<td>0.37</td>
<td>-1.0 ± 2.7t</td>
</tr>
<tr>
<td>Cholesterol (mg)</td>
<td>239.1 ± 91.5</td>
<td>254.8 ± 90.8</td>
<td>0.31</td>
<td>-62.0 ± 68.9t</td>
<td>-22.8 ± 66.4t</td>
<td>0.00</td>
</tr>
<tr>
<td><strong>Anthropometry</strong></td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Body weight (kg)</td>
<td>79.2 ± 14.9</td>
<td>80.3 ± 12.0</td>
<td>0.63</td>
<td>-1.3 ± 1.8t</td>
<td>-0.6 ± 1.9t</td>
<td>0.01</td>
</tr>
<tr>
<td>Body Mass Index (kg/m²)</td>
<td>28.1 ± 4.3</td>
<td>29.2 ± 4.8</td>
<td>0.15</td>
<td>-0.5 ± 0.6t</td>
<td>-0.2 ± 0.7t</td>
<td>0.01</td>
</tr>
<tr>
<td>Waist circumference (cm)</td>
<td>94.3 ± 12.1</td>
<td>97.7 ± 10.3</td>
<td>0.08</td>
<td>-1.6 ± 4.9t</td>
<td>-1.7 ± 5.2t</td>
<td>0.43</td>
</tr>
<tr>
<td>Waist-hip circumference ratio</td>
<td>0.89 ± 0.07</td>
<td>0.90 ± 0.09</td>
<td>0.36</td>
<td>-0.0 ± 0.04</td>
<td>-0.01 ± 0.05t</td>
<td>0.13</td>
</tr>
</tbody>
</table>

1 Joule = 0.24 cal

*Two-sided P values for differences in baseline measures between intervention and control group.

**One-sided P value for difference in change from baseline between intervention and control group.

***P value with multilevel analysis.

†Significant difference in changes after 6 and 12 months compared to baseline within group (one-sided P value < 0.05).
### Organization of consultations

FP consultations: number: 1-3; duration: ± 10 minutes; time in-between: 2 weeks. The intervention stops if no progress is made to the next stage or if the patient relapse to former stage.

Dietician consultations: number: 3; duration: first consultation 30-40 minutes, second and third consultations 10-15 minutes; time in-between: 2 and 8 weeks.

### Classification of stages of change for reduction of dietary fat intake

Before the first FP consultation: an algorithm in combination with the results of fat intake.

At the beginning of the second and third FP consultation: with the help of questions according to the FP manual.

### Content of information packages

A. Feedback on baseline values (dietary intake, serum lipids, anthropometry).

B. Focus on consciousness raising of dietary behavior. Disadvantages mentioned by the patient for eating less fat are discussed and advantages are emphasized by the FP. Education material: test of fat consumption to fill in at home to determine which products make an important contribution to the (saturated) fat intake.

C. Focus on motivation to change dietary behavior. Education material: short food variation list, in which alternatives low in fat and/or high in unsaturated fat are given for products high in saturated fat, to give the patient insight in possible dietary changes.

D. Focus on practical implications to change dietary behavior. Counseling to reduce saturated fat intake and to increase unsaturated fat intake. Advise on reduction of energy and total fat intake to patients high in body weight. For all advice, Dutch dietary guidelines are taken into account. First consultation: living and nutritional habits are asked to enable nutrition advice tailored to the patient's individual situation. This is worked out together with the patient in the form of an example of menus for a whole day. Education materials: brochure about a better fat use and an extensive food variation list. Second consultation: discussion of progress and barriers. Third consultation: discussion of results of second measurements, barriers and expectations.

*Action stage comprises original preparation and action stages.*
change. Preparation and action stages were considered 1 stage (action stage) in this study, given the required nutrition education. Figure 2 summarizes the intervention procedure: 1) counseling aimed at raising consciousness about dietary behavior in the precontemplation stage, 2) motivation to change dietary behavior in the contemplation stage, and 3) if a patient decided to change (action stage), information about practical aspects of dietary change and discussion of referral to a dietician. The intervention was conducted by the patient’s own FP. All patients were referred to the same dietician. Protocols for the FPs and the dietician had been developed and tested prior to the study and were discussed by FPs and the dietician in pre-study group sessions.

**Data analyses**

Differences between groups at baseline and follow-up were tested with unpaired t-tests for continuous variables and with chi-square tests for categorized variables. If the number of observations within 1 cell was less then 5, a Fisher’s exact test was used instead of a chi-square test. Differences within subjects were tested with a paired t-test. *P* values less than 0.05 were considered significant. Because of clustering of patients within practices, a multilevel analysis was also carried out (level 1 patient, level 2 practice). All analyses were performed on the basis of intention to treat. SAS version 6.12 was used for the statistical analyses (SAS Institute Inc., Cary, NC, USA).

**Results**

**Study population**

The study sample was predominantly female (73%), poorly educated (68%), with an average age of 58 years (see Table 1 for definitions of educational level). Of the cardiovascular risk factors, hypertension was present in 92% of participants, type 2 diabetes mellitus in 6%, both disorders in 2%, and a family history of heart disease in 25%. No significant differences were found between the intervention group (n=71) and the control group (n=72) (Table 1). Table 2 demonstrates that the 2 study groups also showed comparable baseline measures according to dietary intake, anthropometry, and serum lipid levels. The mean BMI of the total group of subjects was 28.7 kg/m²; 83% had a BMI higher than 25 and the majority had high total cholesterol and dietary fat intake.

**Intervention-related measures**

At baseline, 51% of the patients in the intervention group were classified in the precontemplation stage, 24% in the contemplation stage, and 25% in the action stage. They had consulted their FP once (n=53) or twice (n=18) before they were referred to the study dietician (n=60). Eleven patients were not referred to the dietician because they did not reach the action stage. All of the referred patients but one received 3 consultations with the dietician. In the control group, 24% of the patients discussed nutrition issues with their FP, 57% read nutrition brochures related to cardiovascular topics, and 1% (7) were referred to a dietician.

**Changes at follow-up measurements**

After 6 months (Table 2) total energy intake was reduced by 1.4 and 0.6 MJ in the intervention and control groups, respectively; total fat intake by 7.9% and 2.2% of total energy, and saturated fat intake by 3.4% and 0.8% of total energy. The reductions were significantly larger in the intervention group, except for unsaturated fat. This was also reflected in risk factors: body weight and BMI declined significantly more in the intervention group (1.5 kg body weight weight) than in the control group (0.6 kg body weight weight). We found no significant differences between groups for waist circumference and waist-hip ratio.

The reduced fat intake in the intervention group was maintained at 12 months, although the differences were smaller. Changes in energy intake and anthropometric values at this time no longer differed significantly with multilevel analysis. During the 12 months of the study, slight reductions were found for serum total cholesterol (intervention group: 2.3 mg/dL, controls: 6.2 mg/dL), LDL cholesterol (intervention group: 6.2 mg/dL, controls: 7.7 mg/dL), and triglycerides (intervention group: 0.8 mg/dL, controls: 3.1 mg/dL). HDL cholesterol increased slightly in both groups (3.9 mg/dL in the intervention group, 2.7 mg/dL in the control group). However, none of these differences were significant. There were no significant changes in smoking or physical activity (*P* values of chi-square tests per measurement moment were >0.85), and none of the patients was prescribed a cholesterol-lowering drug. The significance of the *P* values of the differences in variables between the first and last measurement moment did not change when multilevel analysis was performed, except for body weight (Table 2).

**Discussion**

Nutrition counseling by an FP based on the Stages-of-Change Model, with referral to a dietician in the action stage, successfully changed dietary behavior after 6 months in patients at elevated risk for cardiovascular disease. This success was accompanied by reductions in body weight. Differences in fat intake.
were sustained at 12 months, but this was not reflected in lower serum lipid concentrations. Initial reductions in energy intake and anthropometric values did not persist after 1 year. Our findings are in line with other dietary intervention studies in family practice that report improved dietary habits but no significant effect on objective cardiovascular risk factors such as body weight and blood lipids.29-31 The uniqueness of our study is that it is the first randomized controlled trial in family practice based on the Stages-of-Change Model in which nutrition counseling is managed by the FP.

The reductions we found in total serum cholesterol concentrations (0.9% in the intervention group and 2.3% in the control group after 12 months) were smaller than the 3%-6% suggested by a systematic review of individualized nutrition counseling in free-living subjects.32 The observed reduction in our study is also less than predicted by the Keys equation.33 We do not have a clear explanation for this. It is possible that patients in the intervention group gave more socially desirable answers to the food frequency questionnaire than patients in the control group, due to the more extensive nutrition guidance in the intervention group.

The dropout rate in our study was low: 91% of the patients completed the trial, a notable strength of the study. This may be due to the fact that the participating patients were recruited and treated by their own FP, and may in part account for the small effect size as we avoided the selective participation of those patients who were most motivated for change. The education level of the study sample was low compared with the Dutch population at similar age,34 and this could also have resulted in smaller differences between intervention and control groups. In addition, it has been found that CHD patients who are obese and do not use lipid lowering drugs are less likely to follow recommended cholesterol-lowering diets.35 However, all of these factors make our study representative of the circumstances FPs can encounter patients who are willing to change their food habits. Further, we reached a high percentage of poorly educated people, who are particularly vulnerable.36 We recommend examining whether education materials need to be better aimed at people with a low socio-economic status.

ACKNOWLEDGMENTS - This research was supported by the Netherlands Heart Foundation under grant no. 97.106 and by Bayer. We are grateful to the staff of the NMP family practices and their patients, without whom this study would not have been possible. We extend special thanks to the dieticians José Veen and Els Siebelink and to all the research assistants, especially to Marjolein Homs.

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