Comparison of the effect of individual dietary counselling and of standard nutritional care on weight loss in patients with head and neck cancer undergoing radiotherapy

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and continued during and after cancer treatment, and the implementation may be more successful if a dietitian is involved\(^{(22–24)}\).

Although many studies demonstrate the benefits of full nutritional intervention programme, including dietary counselling, there is little evidence for the potential added value of a professional dietitian’s support. In the current health care system, there is a need to justify resources and to demonstrate the effects of individual dietary counselling.

The present study was designed to evaluate the effect of individual dietary counselling (IDC) by a dietitian for patients with oral cavity, oropharyngeal or hypopharyngeal cancer undergoing radiotherapy before, during and after the treatment. The objective was to investigate whether individualised dietary counselling by a dietitian would better maintain a patient’s body weight, and thus prevent malnutrition compared to standard nutritional care (SC).

**Experimental methods**

**Study design**

A prospective clinical study on the impact of dietary counselling on unintended weight loss, BMI (kg/m\(^2\)) and malnutrition in patients with head and neck carcinomas was carried out between January 2005 and February 2007 at the Radboud University Nijmegen Medical Centre. Patients with primary squamous cell carcinoma in the oral cavity, oropharynx or hypopharynx, age ≥ 18 years, stages II–IV (International Union Against Cancer TNM classification of malignant tumours) were included\(^{(25)}\). Patients were treated depending on stage, location of the tumour and general health conditions by radiotherapy, combined surgery, and radiotherapy or concomitant radio-chemotherapy in accordance with the guidelines of the Dutch Cooperative Head and Neck Oncology Group\(^{(4,5)}\).

From 2005 till 2007, dietary counselling was given to all patients by a dietitian before the start of radiotherapy.

Next, patients were assigned to one of the intervention groups based on their postal code. One group continued to receive IDC by a dietitian throughout the entire treatment and rehabilitation period (IDC). The other group was passed on to trained nurses for SC during radiotherapy and thereafter.

At baseline (the first visit to the outpatient clinic at the departments of Otorhinolaryngology or Oral and Maxillofacial Surgery), patients’ characteristics including age, sex, actual body weight (kg) and height (m) were registered. Parameters studied from baseline through to rehabilitation included body weight (kg), BMI (kg/m\(^2\)); <18·5 = underweight; 18·5–25 = normal range; ≥25 = overweight; ≥30 = obese)\(^{(26)}\) and malnutrition (unintended weight loss ≥5 % in 1 month and/or ≥10 % in 6 months)\(^{(3,27)}\). Body weight was measured on a calibrated one decimal Seca scale (delta nr 707; Seca, Hamburg, Germany) wearing lightweight clothing under the same conditions.

Measurements were taken at five pre-determined time points for each treatment modality (Fig. 1).

**Nutrition policy**

**Individual dietary counselling.** IDC in the present study is defined as individualised and intensive dietary counselling by dietitians focused on maintaining and/or improving a patient’s energy and protein intake. Counselling is based on four nutritional guidelines\(^{(4,5,28,29)}\).

These guidelines describe treatment-related symptoms that influence dietary intake, and offer advice on how to deal with these symptoms, including dietary counselling strategies, when dietary counselling is necessary and when to start tube feeding (nasogastric/gastrostomy). To meet the specific nutritional goals for each patient, individual directions for dietary counselling were given. The patient’s energy requirement was estimated at > 125 kJ to a maximum of 167 kJ (> 30– ≤ 40 kJ) per kg of actual body weight depending on activities, metabolic stress, abnormal losses and treatment. The protein requirement was normally estimated at > 1·0 to a maximum of 1·5 g of protein per kg ideal body weight depending on bedsores or other inflammatory disorders which require extra protein intake. Ideal body weight was estimated as follows: (1) if a patient was underweight (BMI < 20·0 kg/m\(^2\)) ideal weight (kg) = 20·0 kg/m\(^2\) × height (m\(^2\)); (2) if a patient was overweight (BMI > 25·0 kg/m\(^2\)) ideal weight (kg) = 25·0 kg/m\(^2\) × height (m\(^2\)); a patient having a BMI of 20–25 kg/m\(^2\) was considered to be of ideal weight. IDC involved the prescription of a therapeutic diet with regular food which was adjusted to the individual’s usual diet, thereby recognising personal eating patterns, feasible consistency and preferences. The dietitian would also take other relevant factors into consideration, namely the need for alleviation or arrest of local symptoms, as well as psychological factors and digestive and absorptive capacity. The prescription included the type, amount and frequency of feeding, and specified the energetic/protein level that had to be attained. If a patient’s oral intake continued to diminish, energy- and protein-rich oral nutrition supplements were supplied separately or together with normal meals. Supplements (high energetic drinks) were offered to patients in their preferred flavours, and they were instructed to use them as drinks to be consumed between meals or in combination with snacks between meals. When all the above-mentioned methods were insufficient to maintain the required energy and protein intake for patients, nutritional intake was achieved through tube feeding. Gastrostomy tubes were used in cases where tube feeding was expected to be necessary for a

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>Treatment</th>
<th>Early rehabilitation</th>
<th>Rehabilitation</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 (Baseline)</td>
<td>2</td>
<td>3</td>
<td>4</td>
</tr>
</tbody>
</table>

Fig. 1. Data collection: weight in kg (1–5) measured during diagnosis, treatment, early rehabilitation and rehabilitation.
period longer than 6 weeks. From the start of the radiotherapy until at least 2 months after the treatment, patients were checked twice before and at least once a week during and after the radiotherapy. The dietician consultation consisted of twelve to fifteen visits, but it could be arranged more frequently when requested. The dietician was available to answer any queries or provide more information. When necessary, dietician domiciliary care was arranged for the subsequent period.

Standard nutritional care by a nurse. Patients were seen twice by a dietician before radiotherapy (IDC policy) and when radiotherapy started they were passed on to SC by a nurse. This is as follows: body weight of each patient was measured at least once a week on a calibrated scale wearing lightweight clothing.

Nutritional advice was based on a guideline specifically focused on treatment symptoms such as pain, nausea, xerostomia, mucositis and dysphagia, and how to deal with nutrition(29).

Interventions were mostly focused on evaluating pain due to radiation mucositis and adjusting pain medication. When nutritional intake seemed insufficient, advice on high energetic liquid nutrition and/or high kiloenergetic/protein supplements was given. If the above-mentioned methods failed to work, a nasogastric tube was used for patients who had lost more than 10% of their body weight. Gastrostomy tubes were used in cases where tube feeding was expected to be necessary for a period exceeding 6 weeks.

The frequency of nutritional advice was at least once a week, but this increased when more problems occurred (daily, if necessary, depending on the treatment phase and symptoms). Patients were seen at the outpatient clinic or contacted by phone up to a minimum of 3–4 months after the treatment. Trained nurses offered patients support during and after the treatment in all areas concerning illness and side effects such as nutritional, psychological and skin problems, pain and constipation. This support was supervised by a radiation oncologist.

Statistical analysis

Data processing and statistical analysis were performed using Statistical Package for Social Sciences version 16.0 (SPSS, Inc., Chicago, IL, USA).

The percentage of unintended weight loss was calculated using baseline weight as a reference point.

A descriptive statistical analysis was performed for the two groups of patients. Differences in the patients’ characteristics were evaluated by \( \chi^2 \) test. Differences in the mean change in outcome between and within groups were tested by independent and dependent Student’s \( t \) tests, respectively. Fisher’s exact probability test was performed to test the difference in malnutrition between the treatment groups. The level of statistical significance was set at \( P<0.05 \), two-sided.

The present study was conducted according to the guidelines laid down in the Declaration of Helsinki, and all procedures involving human subjects/patients were approved by the Committee on Research Involving Human Subjects of the Radboud University Nijmegen Medical Centre. Written informed consent was obtained from all subjects/patients.

### Table 1. Baseline characteristics of the patients

<table>
<thead>
<tr>
<th>Variable</th>
<th>IDC (n=20)</th>
<th>SE</th>
<th>SC (n=18)</th>
<th>SE</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>63.8</td>
<td>1.9</td>
<td>61.6</td>
<td>2.1</td>
</tr>
<tr>
<td>Height (m)</td>
<td>1.70</td>
<td>0.00</td>
<td>1.71</td>
<td>0.02</td>
</tr>
<tr>
<td>Sex (male)</td>
<td>14.0</td>
<td></td>
<td>8.0</td>
<td></td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>69.3</td>
<td>3.4</td>
<td>71.0</td>
<td>3.8</td>
</tr>
<tr>
<td>BMI (kg/m(^2))</td>
<td>23.9</td>
<td>1.1</td>
<td>24.0</td>
<td>0.9</td>
</tr>
<tr>
<td>(&lt;18.5)</td>
<td>3.0</td>
<td></td>
<td>1.0</td>
<td></td>
</tr>
<tr>
<td>18.5–25.0</td>
<td>10.0</td>
<td></td>
<td>12.0</td>
<td></td>
</tr>
<tr>
<td>(&gt;25.0)</td>
<td>7.0</td>
<td></td>
<td>5.0</td>
<td></td>
</tr>
<tr>
<td>T stage</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>T2</td>
<td>6.0</td>
<td></td>
<td>11.0</td>
<td></td>
</tr>
<tr>
<td>T3</td>
<td>7.0</td>
<td></td>
<td>4.0</td>
<td></td>
</tr>
<tr>
<td>T4</td>
<td>7.0</td>
<td></td>
<td>3.0</td>
<td></td>
</tr>
<tr>
<td>N stage</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>N⁰</td>
<td>9.0</td>
<td></td>
<td>5.0</td>
<td></td>
</tr>
<tr>
<td>N ≥ 1</td>
<td>11.0</td>
<td></td>
<td>13.0</td>
<td></td>
</tr>
<tr>
<td>Tumour site</td>
<td>10.0</td>
<td></td>
<td>3.0</td>
<td></td>
</tr>
<tr>
<td>Oral cavity</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Oropharynx</td>
<td>7.0</td>
<td></td>
<td>12.0</td>
<td></td>
</tr>
<tr>
<td>Hypopharynx</td>
<td>3.0</td>
<td></td>
<td>3.0</td>
<td></td>
</tr>
<tr>
<td>Treatment</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Radiotherapy</td>
<td>6.0</td>
<td></td>
<td>7.0</td>
<td></td>
</tr>
<tr>
<td>Chemo/radiotherapy</td>
<td>6.0</td>
<td></td>
<td>5.0</td>
<td></td>
</tr>
<tr>
<td>Surgery/radiotherapy</td>
<td>8.0</td>
<td></td>
<td>6.0</td>
<td></td>
</tr>
</tbody>
</table>

IDC, individual dietary counselling; SC, standard nutritional care by a nurse.

* BMI (kg/m\(^2\)): <18.5, underweight; 18.5–25.0, healthy weight; >25.0, overweight (WHO)(26).

### Results

A total of thirty-eight patients met the inclusion criteria. The IDC group included twenty patients, and the SC group consisted of eighteen patients. Table 1 summarises baseline characteristics of the two groups. There were no significant differences between age, height, sex, weight, BMI, T stage, N stage, tumour site and treatment.

### Percentage of unintended weight loss

The percentage weight changes for the two groups are shown in Fig. 2.
The IDC group had a maximum of 3 % unintended weight loss 2 weeks after the treatment. This was approximately the same for the SC group. Two months after the treatment, the IDC group started to gain weight (1 % weight gain), while the SC group continued to lose 1·5 % weight. Two months after the treatment, weight loss was significantly different between SC and IDC groups (P=0·03).

**BMI**

Most of the patients in IDC and SC groups had a normal BMI (18·5 kg/m² ≤ BMI < 25 kg/m²) throughout the study (Fig. 3). No patients were underweight (BMI < 18·5 kg/m²). BMI for the IDC and SC groups did not differ significantly at any of the time points.

**Malnutrition**

Table 2 displays the prevalence of malnutrition from diagnosis until 2 months after the treatment. Before treatment, four of twenty patients in the IDC group and three of eighteen patients in the SC group were malnourished. During the treatment period, the IDC and SC groups remained stable. Two weeks after the treatment, there was a significant difference in malnutrition between the IDC (0/20) and SC (5/18) groups (P=0·02).

Two months after the treatment, the highest prevalence of malnutrition was seen for the SC group (3/18).

The prevalence of malnutrition by T stages is shown in Table 3. No significant differences were found between the T stages.

**Discussion**

Malnutrition, defined as ≥ 5 % of unintended weight loss/month or ≥ 10 % of unintended weight loss/6 months, is the most commonly used parameter in clinical practice for head and neck cancer[3,27]. In the literature, the prevalence of malnutrition is associated with higher mortality and morbidity rates, shorter failure-free survival and poorer quality of life among radiotherapy patients[11–17]. It is therefore essential to control unintended weight loss/malnutrition and maintain an efficient nutritional status for patients. The present study demonstrates the beneficial effect of IDC on weight change, BMI and malnutrition for patients with oral cavity, oropharyngeal and hypopharyngeal cancers treated with radiotherapy compared to SC.

Two different groups (IDC and SC) were compared in the present study. Patients’ characteristics were evenly distributed. Treatment-induced side effects of patients receiving radiotherapy to the head and neck area peaked at the end of the treatment course and continued for 2 or more weeks after the treatment(20). This is reflected by a sharp weight reduction during this period, which continued until 2 months after the treatment, specifically in the SC group, while the IDC group started gaining weight 2 weeks after the radiotherapy. There was a significant difference in unintended weight loss 2 months after the treatment for the SC group compared with the IDC group.

The IDC group started with the same proportion of patients with malnutrition (4/20) vs SC (3/18) at diagnosis where malnutrition in the SC group increased over time. A significant difference in the prevalence of malnutrition for the early rehabilitation period was seen between the groups (P=0·02), which affirms that dietary counselling recovers malnutrition where SC increases malnutrition.

While looking at the overall results of the present study, the most striking differences are observed in the (early) rehabilitation period. Two months after the treatment, weight and BMI in the SC group were still declining. In future research, a longer follow-up is therefore required to determine the nadir of weight loss and the duration of full recovery. Also, more data about nutritional status should be included such as a comprehensive nutritional assessment tool (e.g. subjective global assessment or patient-generated subjective global assessment), anthropometric measurements (e.g. bioimpedance and/or handgrip) and food intake (e.g. dietary history or FFQ). Besides, it should be useful to include other endpoints such

**Table 2.** Prevalence of malnutrition* in individual dietary counselling (IDC) and standard nutritional care by a nurse (SC) groups from diagnosis until rehabilitation

<table>
<thead>
<tr>
<th>Interval</th>
<th>Number of patients per nutrition intervention</th>
</tr>
</thead>
<tbody>
<tr>
<td>Diagnosis</td>
<td>ICD (n=20)</td>
</tr>
<tr>
<td>Treatment</td>
<td>4·0</td>
</tr>
<tr>
<td>Early rehabilitation</td>
<td>0·0†</td>
</tr>
<tr>
<td>Rehabilitation</td>
<td>1·0</td>
</tr>
</tbody>
</table>

* Malnutrition was defined as ‘unintended weight loss ≥ 5 % within 1 month’.
† Prevalence of malnutrition in IDC and SC groups was significantly different (P=0·02).

**Table 3.** Prevalence of malnutrition* per T stage from diagnosis until rehabilitation

<table>
<thead>
<tr>
<th>Interval</th>
<th>Number of patients per T stage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Diagnosis</td>
<td>T2 (n=17)</td>
</tr>
<tr>
<td>Treatment</td>
<td>2·0</td>
</tr>
<tr>
<td>Early rehabilitation</td>
<td>5·0</td>
</tr>
<tr>
<td>Rehabilitation</td>
<td>3·0</td>
</tr>
</tbody>
</table>

* Malnutrition was defined as ‘unintended weight loss ≥ 5 % within 1 month’.
as quality of life, mortality, response to treatment and length of hospital stay.

These results indicate that IDC significantly contributes to maintaining and improving weight loss and malnutrition of head and neck cancer patients undergoing radiotherapy. This is an important clinically relevant finding which confirms the research objective of the present study.

A randomised study performed by Ravasco et al. produced similar results. The study demonstrated that individualised dietary counselling (based on regular foods) for head and neck cancer patients undergoing radiotherapy is the most effective way of improving patients’ nutritional intake, nutritional status and quality of life by the end of the treatment until 3 months after the treatment.

However, Ravasco used Ottery’s Subjective Global Assessment (unscored patient-generated subjective global assessment) to determine malnutrition. In this system, unintended weight loss is one of the deciding factors besides symptoms, metabolic stress and subcutaneous fat.

A randomised controlled trial by Isenring et al. showed that dietary counselling using the American Dietetic Association – Medical Nutrition Therapy radiation oncology protocol resulted in a significant reduction in unintended weight loss from the start of radiotherapy until 3 months after the treatment compared with their standard practice (which consisted of general nutritional advice by a nurse and a booklet). Furthermore, the present study revealed an improvement in nutritional status (patient-generated subjective global assessment), dietary intake and quality of life in the dietary counselling group when compared with the standard practice. This is the first prospective randomised controlled trial. A limitation of the present study is that two different tumour locations were involved (gastrointestinal and head and neck cancers).

Dawson et al. reported a significant reduction in unintended weight loss in oral cancer patients undergoing combined modality treatment (surgery + radiotherapy) with increased dietary supervision compared with standard dietary counselling after revising their dietary protocol. However, this was not compared to nutritional standard care without dietary counselling. It is interesting to note that while there have been only a handful of studies in this area by different research teams, they all have demonstrated benefits of dietary counselling like the present study.

To summarise, early and intensive IDC produces clinically relevant effects in terms of decreasing unintended weight loss and malnutrition compared with SC in patients with head and neck cancer undergoing radiotherapy. This is especially the case early after the treatment and during the rehabilitation period.

The guideline used by the Radboud University Nijmegen Medical Centre Department of Dietetics for head and neck cancer patients is a useful guide in supplying the level of dietary support required. In clinical practice, patients should receive regular and individualised dietary counselling from diagnosis until at least 2 months after radiotherapy and probably longer.

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References


