Pre-cachexia and cachexia at diagnosis of stage III non-small-cell lung carcinoma: an exploratory study comparing two consensus-based frameworks

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Abstract
Despite the development of consensus-based frameworks to define cancer cachexia, the validity and usefulness of these frameworks are relatively unknown. The aim of the present study was to study the presence of pre-cachexia and cachexia in patients with stage III non-small-cell lung carcinoma (NSCLC) by using a cancer-specific framework and a general framework for cachexia, and to explore the prognostic value of pre-cachexia and cachexia. In forty patients at diagnosis of stage III NSCLC, weight loss, fat-free mass, handgrip strength, anorexia and serum biochemistry, assessed before the first chemotherapy, were used to define ‘cancer cachexia’ or ‘cachexia’. The cancer-specific framework also classified for pre-cachexia and refractory cachexia. Additionally, quality of life was assessed by the European Organisation for Research and Treatment of Cancer – Quality of Life Questionnaire C30. Groups were compared using independent t tests, ANOVA, Kaplan–Meier and Cox survival analyses. Based on the cancer-specific framework, pre-cachexia was present in nine patients (23 %) and cancer cachexia was present in seven patients (18 %). Cancer cachexia was associated with a reduced quality of life (P=0.03) and shorter survival (hazard ratio (HR) = 2.9; P=0.04). When using the general framework, cachexia was present in eleven patients (28 %), and was associated with a reduced quality of life (P=0.08) and shorter survival (HR = 4.4; P=0.001). In conclusion, pre-cachexia and cachexia are prevalent in this small population of patients at diagnosis of stage III NSCLC. For both frameworks, cachexia appears to be associated with a reduced quality of life and shorter survival. Further studies are warranted to more extensively explore the validity and prognostic value of these new frameworks in cancer patients.

Key words: Pre-cachexia: Cachexia: Non-small-cell lung cancer

Cachexia is a complex metabolic syndrome characterised by ongoing loss of body weight and skeletal muscle mass, which cannot be fully reversed by conventional nutritional support11. The pathophysiology of cachexia encompasses a negative protein and energy balance, driven by a variable combination of reduced food intake and abnormal metabolism. Cachexia is frequently observed in patients with cancer, and is associated with progressive functional impairment, intolerance to anticancer treatment and shorter survival11–31. The severity of cachexia in patients with cancer varies from non-symptomatic inflammatory derangements and minimal weight and muscle loss in the early stage to severe muscle wasting and low performance status in patients not responding to anticancer treatment44.

In order to define and stage cachexia, a number of frameworks in patients with chronic diseases1,5 and cancer4,6–8 have been described. Recently, an international expert group proposed a conceptual framework for cancer cachexia, with a classification for three stages of clinical relevance: pre-cachexia, cachexia and refractory cachexia44. Overall, existing instruments use slightly different nutritional and inflammatory parameters and cut-off points to define pre-cachexia and cachexia.

Despite the growing understanding of the pathophysiology and staging of cachexia, assessment of cachexia in clinical

Abbreviations: CRP, C-reactive protein; EORTC-QLQC30, European Organisation for Research and Treatment of Cancer – Quality of Life Questionnaire C30; ESPEN, European Society for Parenteral and Enteral Nutrition; FFM, fat-free mass; HR, hazard ratio; NSCLC, non-small-cell lung cancer; REE, resting energy expenditure; TEE, total energy expenditure; VAS, visual analogue scale.

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practice is limited. These studies clearly showed the occurrence of weight loss and features of cachexia in patients with cancer.

In patients with lung cancer, high prevalences of involuntary weight loss have been reported\(^9\text{--}^{11}\). Lung cancer is frequently associated with cachexia. Weight loss in patients with lung cancer was associated with systemic inflammation, loss of muscle mass, an increased acute-phase response, decreased levels of the anabolic hormone insulin-like growth factor-I\(^\text{12}\) and hypermetabolism\(^\text{12,13}\). Weight loss was also associated with reduced quality of life\(^\text{14}\), response to chemotherapy\(^\text{15}\) and survival\(^\text{9,16}\) in patients with lung cancer.

The staging of cachexia in patients with lung cancer has not been described, but could help clinicians to decide on early interventions or cachexia treatment. Up to now, the validity and usefulness of cachexia instruments in patients with cancer is unknown, and the recognition and nutritional management of cancer cachexia remains unsatisfactory\(^\text{1}\).

Comprehensive data of cancer populations could give more insight into the pathophysiology of (pre)cachexia, and could be used to apply cachexia frameworks and to investigate the outcomes and differences between frameworks. Therefore, we aimed to retrospectively study the presence of (pre)cachexia at diagnosis of stage III non-small-cell lung cancer (NSCLC), using recently described consensus-based frameworks\(^\text{1,4,5}\), and to explore the prognostic value of pre-cachexia and cachexia. Second, we explored quality of life, and nutritional and inflammatory parameters associated with (pre)cachexia. We hypothesise that (pre)cachexia is present in this locally advanced patient population and that cachexia is associated with a decreased quality of life and shorter survival.

Materials and methods

Patients

Between March 2005 and October 2007, forty patients with histologically or cytologically proven stage III NSCLC, aged 18–80 years and having a life expectancy of at least 3 months, were included at the start of concurrent chemoradiotherapy. Patients were excluded if they had undergone surgery, chemotherapy or radiotherapy during the previous month; if they had oedema, ascites or severe co-morbidities; or if they used high-dose corticosteroids or fish oil supplements.

Data used for the present retrospective analysis were collected at the inclusion for a prospective double-blind randomised controlled trial that has been carried out at our centre from 2005 to 2008. Out of fifty-five enrolled patients, four patients did not meet the inclusion criteria, nine patients refused to participate and two had disease progression. Thirty patients did not meet the inclusion criteria, nine patients refused to participate and two had disease progression (Fig. S1, available online). We used the baseline and survival data of forty patients, irrespective of the intervention in the trial. After carrying out baseline measurements, patients were randomly assigned to receive two cans per d of either a protein- and energy-dense oral nutritional supplement containing n-3 PUFA or an isoenergetic control oral nutritional supplement during 5 weeks of chemoradiotherapy\(^\text{17}\).

Throughout chemoradiotherapy, the diétitian monitored dietary intake and provided dietary counselling. Tube feeding was indicated in the case of an (expected) oral intake of <75% of energy requirements for more than 3 d, combined with the inability to increase energy intake by oral food or sip feeds.

The present study was conducted according to the guidelines laid down in the Declaration of Helsinki, and all procedures involving human patients were approved by the Medical Ethics Committee of the VU University Medical Center, Amsterdam, The Netherlands. Written informed consent was obtained from all patients.

Baseline measurements

At baseline, before the start of chemoradiotherapy, weight loss, BMI, fat-free mass (FFM), energy expenditure, anorexia, inflammation, muscle strength, quality of life and physical activity were assessed.

Weight loss and BMI

Pre-illness weight, unintentional weight loss in the last month and during the last 6 months and height were recorded. Body weight, without shoes and wearing light clothing, was measured on a compact digital flat scale (SECA 888) to the nearest 0·2 kg. BMI was calculated by dividing body weight (kg) by the square of the height (m).

Fat-free mass

Bioelectrical impedance spectroscopy (Hydra 4200, Xitron Technologies) was performed to assess FFM. Whole-body resistance was measured with four surface electrodes placed on the right wrist and ankle, as previously described\(^\text{18}\). Briefly, the principle was based on the application of a variable electrical current between 50 and 700 μA produced by a generator and applied to the skin using adhesive electrodes (3M red Dot Ag/AgCl) with the subject lying supine\(^\text{19}\). FFM was calculated from resistance and reactance at the frequency of capacitance by using the Kyle Geneva equation\(^\text{20}\).

The phase angle of bioelectrical impedance at 50kHz was calculated using the following equation: phase angle = (resistance/reactance) \times (180/\pi). The cut-off point for patients with lung cancer, described by Gupta et al.\(^\text{21}\), was used to classify patients with a low (≤5·3) and high (>5·3) phase angle.

Energy expenditure

Resting energy expenditure (REE) was measured by a ventilated hood system (Deltatrac, Datex); CO\(_2\) production (VCO\(_2\)) and O\(_2\) consumption (VO\(_2\)) were measured at complete rest for a period of 30 min. REE was calculated using a modified Weir equation\(^\text{22,23}\). To estimate total energy expenditure (TEE), 30% was added to REE, assuming a physical activity level of 1·3 for sedentary patients with cancer\(^\text{24}\).

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Patients recorded their appetite on a visual analogue scale (VAS), 10 cm in length.\(^{[25]}\). Patients’ energy intake, assessed by a 24 h dietary recall, was expressed as percentage of TEE. Anorexia and/or reduced food intake were identified by the presence of appetite < 5 cm (VAS), energy intake < 84 kJ/kg body weight per d (84 kJ (20 kcal)/kg)\(^{[11]}\) or energy intake < 70 % of TEE\(^{[1]}\).

**Inflammation.** Non-fasting blood samples were taken simultaneously with usual blood samples for chemotherapy. Plasma concentrations of C-reactive protein (CRP) were measured with an automated latex-enhanced immunoturbidimetric assay on a Modular P analyser (reference: 0–8 mg/l)\(^{[26]}\). Serum IL-6 was measured by commercially available ELISA (Pelikine compact human ELISA kits, Sanquin) (reference: 0–4 pg/ml). Whole-blood Hb was determined by spectrophotometry on a Cell-Dyn Sapphire analyser (Abbott Diagnostics) (reference: ≥ 7–3 mmol/l or ≥ 117 g/l)\(^{[27]}\). Serum albumin concentrations were chemically determined on a Modular P analyser (ACN 760, 11815148 216, Roche Diagnostics) (reference: ≥ 320 g/l)\(^{[28]}\).

**Muscle strength.** Muscle strength was measured by handgrip strength in the non-dominant hand using a hydraulic hand dynamometer (Baseline, Fabrication Enterprises). The patient performed two maximal isometric contractions while the handgrip dynamometer was firmly held between the palms and the index fingers. The average of two measurements was recorded, and elbow flexed at 90°.

**Definition of pre-cachexia and cachexia**

We used two consensus-based frameworks to define cachexia: a cancer-specific and a non-disease-specific general framework. With the cancer-specific framework, we defined pre-cachexia, cancer cachexia and refractory cancer cachexia, as proposed by, respectively, the European Society for Parenteral and Enteral Nutrition (ESPEN) Special Interest Group ‘cachexia–anorexia in chronic wasting diseases’\(^{[3]}\) and an international panel of experts in clinical cancer cachexia research\(^{[4]}\). Second, we used the general framework for cachexia in chronic illness, as described by Evans et al.\(^{[5]}\).

Some of the parameters and cut-off points not specifically given a reference citation in the sections below were not described in frameworks, and therefore retrieved from available unspecified literature and, where necessary, from experts.

**Cancer-specific framework for cachexia**

**Cancer pre-cachexia**\(^{[1]}\):\(^{[1]}\)

1. Unintentional weight loss of 0 to ≤ 5 % during the previous 6 months.
2. Anorexia (the presence of either: appetite < 5 cm (VAS), energy intake < 84 kJ/kg body weight per d (84 kJ (20 kcal)/kg)\(^{[11]}\) or energy intake < 70 % of TEE\(^{[1]}\)).
3. Systemic inflammation (CRP ≥ 8 mg/l, the upper limit of normality).

**Cancer cachexia**\(^{[4]}\):\(^{[4]}\)

1. Weight loss > 5 % during the previous 6 months or BMI < 20 kg/m\(^2\) and weight loss > 2 % or sarcopenia (FFM index < 5th percentile of age- and sex-specific reference values\(^{[34]}\) and weight loss > 2 %).
2. Reduced food intake (the presence of either: appetite < 5 cm (VAS), energy intake < 84 kJ/kg body weight per d (84 kJ (20 kcal)/kg)\(^{[11]}\) or energy intake < 70 % of TEE\(^{[1]}\)).
3. Systemic inflammation (CRP ≥ 8 mg/l, the upper limit of normality)

**Refractory cancer cachexia**\(^{[4]}\):\(^{[4]}\)

1. Variable degree of ‘cancer cachexia’.
2. Cancer disease both pro-catabolic and not responsive to anticancer treatment.
3. Low performance score (Karnofsky Performance Score < 50, indicating that a patient is unable to care for self).
4. < 3 months expected survival.

**General framework**

The non-disease-specific general framework for cachexia\(^{[5]}\) includes the combination of weight loss of ≥ 5 % in 6 months or BMI < 20 kg/m\(^2\), combined with at least three of the following five criteria:

1. Decreased muscle strength.
2. Handgrip strength below the lowest tertile extracted from age- and sex-specific reference values\(^{[29]}\).
(2) Fatigue (score of 3 or 4 according to the EORTC-QLQ-C30 symptom scale\(^{(12)}\)).

(3) Anorexia (the presence of: appetite < 5 cm (VAS), energy intake < 84 kJ/kg body weight per day (84 kcal/kg)\(^{(11)}\) or energy intake < 70% of TEE\(^{(13)}\)).

(4) FFM index below the 10th percentile by age- and sex-specific reference values\(^{(13)}\).

(5) One or more abnormal serum biochemistry parameters: CRP > 5 mg/l, Hb < 120 g/l or 117 g/l, serum albumin < 32 g/l or 30 g/l, 117 g/l, serum albumin < 32 g/l or 30 g/l, IL-6 > 4 pg/ml\(^{(6)}\).

**Statistical analysis**

Statistical analysis was performed using SPSS for Windows (version 17.0, SPSS, Inc.). Groups with no cachexia, pre-cachexia and cachexia were compared for serum biochemistry, REE, quality of life and physical role, emotional, cognitive and social functioning. Independent samples \(t\) tests were performed to compare groups with no cachexia and cachexia. For variables that were not normally distributed, non-parametric tests were performed to compare group differences. Frequencies within groups for nominal characteristics were compared by Pearson’s \(\chi^2\) tests. Differences between three groups (no cachexia, pre-cachexia and cachexia) were tested by one-way ANOVA. Correlations between variables were investigated by Pearson’s correlation tests.

Group survival, from the date of the start of concurrent chemotherapy (from 15 March 2005 until 30 October 2007) until death or follow-up visit (17 November 2011), was generated by the method of Kaplan and Meier and compared by means of the log-rank test. Second, the multivariate Cox’s proportional hazards model was used to analyse means of the log-rank test. Cachexia was the independent variable and the model was adjusted for confounding factor(s) (based on a > 10% change of OR, after adding a single factor: sex, age and/or tumour stage: IIa vs. IIIb). Median survival was displayed with the standard error; \(P\) values < 0.05 were considered to be statistically significant.

**Results**

**Patients**

A total of forty patients with histologically or cytologically proven stage IIa \((n = 16)\) or stage IIIb \((n = 24)\) NSCLC were studied, nineteen females and twenty-one males, with a median age of 57 (range 39–80) years. The average amount of weight loss during the previous 6 months was 1.9 (SD 6.5)% of pre-illness weight. The overall median survival was 25.0 (SD 8.7) months. Baseline patient characteristics are displayed in Table 1.

**Cancer-specific framework**

Using the two consensus-based frameworks of the ESPEN Special Interest Group\(^{(3)}\) and Fearon et al.'s\(^{(4)}\), we classified pre-cachexia in nine patients (23%) and cachexia in seven patients (18%). The remaining twenty-four patients were classified as no-cachexia patients (Table 2). None of the patients met the criteria of refractory cancer cachexia: measurements were carried out at diagnosis, just before starting anticancer treatment, and the Karnofsky performance score was relatively high (70–100) and the expected survival was at least 3 months in all patients. Quality of life was significantly different among no-cachexia, pre-cachexia and cachexia groups (\(P < 0.05\)), but other function scales (such as physical function) did not significantly differ between groups. Survival was non-significantly different between no-cachexia, pre-cachexia and cachexia groups in univariate analysis (24 (SD 11.6) vs. 32 (SD 1.5) v. 9 (SD 9.2) months,

**Table 1. Baseline characteristics for patients with stage III non-small-cell lung cancer, specified for groups with no cachexia, pre-cachexia and cachexia, as defined by different consensus-based frameworks**

<table>
<thead>
<tr>
<th>Overall (n 40)</th>
<th>No cachexia (n 24)</th>
<th>Pre-cachexia (n 9)</th>
<th>Cachexia (n 7)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>57.8 10.1</td>
<td>57.7 11.1</td>
<td>57.0 8.1</td>
<td>59.0 10.0</td>
</tr>
<tr>
<td>Female</td>
<td>0.93*</td>
<td>0.59†</td>
<td></td>
<td></td>
</tr>
<tr>
<td>n</td>
<td>19</td>
<td>12</td>
<td>3</td>
<td>4</td>
</tr>
<tr>
<td>%</td>
<td>47.5</td>
<td>50</td>
<td>33</td>
<td>57</td>
</tr>
<tr>
<td>Tumour stage</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>IIa</td>
<td>0.64†</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>n</td>
<td>16</td>
<td>11</td>
<td>3</td>
<td>2</td>
</tr>
<tr>
<td>%</td>
<td>40</td>
<td>46</td>
<td>33</td>
<td>29</td>
</tr>
<tr>
<td>IIIb</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>n</td>
<td>24</td>
<td>13</td>
<td>6</td>
<td>5</td>
</tr>
<tr>
<td>%</td>
<td>60</td>
<td>54</td>
<td>56</td>
<td>71</td>
</tr>
<tr>
<td>Weight change in previous 6 months (%)</td>
<td>1.3 4.5</td>
<td>0.3 5.5</td>
<td>-0.5 1.0</td>
<td>-11.6 5.0</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>23.9 3.5</td>
<td>24.5 3.5</td>
<td>23.5 2.8</td>
<td>22.5 4.2</td>
</tr>
</tbody>
</table>

* ANOVA (comparing no-cachexia, pre-cachexia and cachexia groups).
† Independent samples \(t\) test for equality of means (comparing no-cachexia and cachexia groups).
‡ Pearson’s \(\chi^2\) test (comparing no-cachexia and (pre)cachexia groups).
Table 2. Number of patients with stage III non-small-cell lung cancer classified as having no cachexia, pre-cachexia and cachexia

<table>
<thead>
<tr>
<th></th>
<th>No cachexia</th>
<th>Pre-cachexia</th>
<th>Cachexia</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>General</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>No cachexia</td>
<td>20</td>
<td>9</td>
<td>0</td>
<td>29</td>
</tr>
<tr>
<td>Cachexia</td>
<td>4</td>
<td>0</td>
<td>7</td>
<td>11</td>
</tr>
<tr>
<td>Total</td>
<td>24</td>
<td>9</td>
<td>7</td>
<td>40</td>
</tr>
</tbody>
</table>

respectively; $P=0.21$ (Fig. 1). Multivariate analysis with no cachexia as the reference category, corrected for sex and tumour stage, showed a significantly shorter survival in patients with cancer cachexia (HR 2.93; 95 % CI 1.03, 8.34; $P=0.04$), but not in patients with pre-cachexia (HR 0.78; 95 % CI 0.30, 2.03; $P=0.62$).

**General framework**

Using the general framework to define cachexia, we identified eleven (28 %) out of forty patients with cachexia and twenty-nine patients (72 %) as having no cachexia (Table 2). The four patients who were classified as cachectic using the general definition, but not when using the cancer-specific framework, did not experience anorexia, but scored positive in at least three other features of the general definition (Table 3). Cachexia tended to be associated with a trend for a lower quality of life ($P=0.08$). Between the general no-cachexia and the cachexia groups, median survival was significantly different (respectively, 32.0 (SD 4.5) v. 10.0 (SD 3.7) months; $P<0.01$) (Fig. 2). In multivariate analysis, corrected for confounding by sex and tumour stage, cachexia remained significantly associated with a shorter survival (HR 4.2; 95 % CI 1.7, 10.0; $P=0.001$).

**Cachexia features**

Approximately 50 % of non-cachectic patients scored positively on cachexia features, such as fatigue, anorexia, reduced handgrip strength and upper arm circumference, or increased CRP. In general, low percentages of patients scored positively on a reduced FFM index, albumin or Hb (Table 3).

For all instruments, groups with cachexia showed higher levels of CRP and IL-6 and a lower Hb and serum albumin than patients with no cachexia ($P<0.01$) (Table 4). CRP was positively correlated with IL-6 ($r=0.55$, $P<0.01$), and negatively correlated with Hb ($r=-0.47$, $P<0.01$) and serum albumin ($r=-0.71$, $P<0.01$). The remaining inflammatory parameters were also significantly correlated with one another. Using different cut-off points for CRP (>5 or 10 mg/l instead of >8 mg/l) did not change the presence of pre-cachexia and cachexia in individual patients (data not shown).

Of all patients, twelve (30 %) had a weight loss of at least 5 % in the previous 12 months or less, three (8 %) had a FFM index below the 5th percentile of reference values, twenty-seven (68 %) had decreased handgrip strength (below the lowest tertile of reference values), nineteen (48 %) experienced fatigue and twenty-three (58 %) experienced anorexia or reduced food intake. When comparing individual levels of inflammatory parameters with their reference values, CRP and serum IL-6 were elevated in, respectively, twenty-eight (70 %) and twenty (50 %) patients, and Hb and serum albumin were decreased in eight (20 %) and seven (18 %) patients, respectively (Table 3).

**Additional parameters**

REE per kg FFM, physical activity and phase angle were non-significantly different between groups. However, physical activity appeared to be lower in cachexia patients (Table 4).

**Discussion**

The purpose of the present explorative study was to study the presence of pre-cachexia and cachexia in patients with stage III NSCLC, by using consensus-based conceptual frameworks, which have not yet been applied or validated in populations of patients with cancer. Second, we explored the association of (pre)cachexia with survival and quality of life. Although we are gaining knowledge on the pathophysiology and treatment of cancer cachexia, little is known about the typical profile and staging of cachexia. We chose to apply the only two available consensus-based frameworks to define cachexia. These frameworks were both comprehensive, but differed in the kind of parameters to define cachexia. The cut-off point of essential parameters, e.g. weight loss, is still a subject of debate. Therefore, we were interested in the outcomes of these two instruments when applied in a small, heterogeneous population of patients with locally advanced cancer.

These frameworks defined cachexia and described the clinical features associated with cachexia. More recently published proposals that aimed to grade the severity of cachexia led to the definition of pre-cachexia. Using these proposals, it is possible to identify cancer patients with pre-cachexia: early-stage cachexia, characterised by moderate systemic inflammation and metabolic alterations, and minimal weight loss.
Patients with pre-cachexia are not always recognised by clinicians or nutritional screening instruments, while nutritional support is expected to prevent progressive loss of body weight and FFM. On the contrary, treatment options for cachexia are limited. In the present population of patients at diagnosis of stage III NSCLC, pre-cachexia was prevalent in 23 %, but only the framework proposed by the ESPEN Special Interest Group defines cachexia (respectively, 18 and 28 % by the cancer-specific and non-disease-specific general framework). Cachexia was also prevalent in the pre-sent population, but the cancer-specific framework and the general framework for cachexia found a different number of patients with cachexia (respectively, 18 and 28 % by the cancer-specific and non-disease-specific general framework).

A number of studies showed the association between survival and weight loss in general cancer populations and in patients with gastrointestinal and lung cancer. One of the first papers on this topic found the combination of weight loss, food intake and systemic inflammation to be related to poor outcome in pancreatic cancer patients. Because the definition of cachexia includes the presence of severe weight loss, the association with survival in the present study is consistent with these findings. The difficulty is that, in the literature, weight loss and cachexia are used disorderly, and that it is not possible to isolate starvation from cancer cachexia. Another component of the cachexia definition is inflammation. Systemic inflammation, amongst others reflected by elevated CRP and hypoalbuminemia, is also negatively associated with survival.

After carrying out baseline measurements at diagnosis, patients received different anticancer treatments and participated in a placebo-controlled randomised controlled trial comparing oral nutritional supplements containing n-3 PUFA with an isoenergetic placebo. Yet, the percentages of patients with (pre)cachexia and survival did not significantly differ among groups with different cancer treatments (data not shown). Preclinical studies suggest that an increased intake of n-3 PUFA decreases the risk of cancer development and...
progression. A few clinical studies support the potential benefit of n-3 PUFA on chemotherapy efficacy or cancer cell proliferation. In the present population, patients who received oral nutritional supplements containing n-3 PUFA did not show a significantly different presence of cachexia or survival than control patients (data not shown).

On average, the present population of patients with stage III NSCLC showed a moderate amount of weight loss (on average 1.9% of pre-illness weight) during the previous 6 months and, consequently, a low prevalence of malnutrition (20%). Other studies in patients with lung cancer (all types and stages) reported high percentages of malnutrition, i.e. 15-30% and 50-61%. A study by Bozzetti & Mariani showed an average weight loss of 9.5% in outpatient with lung cancer. Because we used bioelectrical impedance spectroscopy to assess FFM (and not the ‘gold standard’ dual-energy X-ray absorptiometry), this may have resulted in over-estimation or under-estimation of FFM.

We also showed that approximately 50% of non-cachectic patients scored positively on cachexia features, such as moderate weight loss, systemic inflammation, fatigue, anorexia, reduced handgrip strength and upper arm circumference. The frameworks that we used define patients as pre-cachectic or cachetic when they experience a combination of cachexia features, inflammation and weight loss, which is consistent with the existing knowledge on the pathophysiology of cachexia. The present study used the proposal of the SCReening the Nutritional Status in Oncology (SCRINIO) working group.

**Table 4. Differences in biochemistry, phase angle, resting energy expenditure (REE), physical activity and quality of life between cachexia groups with stage III non-small-cell lung cancer**

(Mean values and standard deviations)

<table>
<thead>
<tr>
<th>Cancer</th>
<th>General</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No cachexia</td>
</tr>
<tr>
<td></td>
<td>(n 24)</td>
</tr>
<tr>
<td>Hb (g/l)</td>
<td>133 ± 13</td>
</tr>
<tr>
<td>Serum albumin (g/l)</td>
<td>383 ± 38</td>
</tr>
<tr>
<td>CRP (mg/l)</td>
<td>30.4 ± 53.6</td>
</tr>
<tr>
<td>Serum IL-6 (pg/ml)</td>
<td>3.2 ± 2.5</td>
</tr>
<tr>
<td>Phase angle (°)</td>
<td>6.9 ± 2.2</td>
</tr>
<tr>
<td>REE (kJ)</td>
<td>6406 ± 895</td>
</tr>
<tr>
<td>REE (kJ/kg FFM)</td>
<td>129 ± 24</td>
</tr>
</tbody>
</table>

* ANDA (comparing no-cachexia, pre-cachexia and cachexia groups). † Independent samples t-test for equality of means (comparing no-cachexia and cachexia groups).

†1k J = 0.239 kcal.

CRP, C-reactive protein; FFM, fat-free mass; PAM, Physical Activity Monitor; EORTC-QLQ-C30, European Organisation for Research and Treatment of Cancer – Quality of Life Questionnaire C30.

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of life parameters, such as physical function. In the literature, an association among nutritional status, inflammation and well-being in lung cancer has been described, but these studies did not assess cachexia in the way we did\(^{45,46}\).

Physical activity is an important indicator of quality of life and performance status in cancer patients\(^{47}\), and found to be reduced in patients with SCLC\(^{48}\) and pancreatic cancer\(^{49}\). The present patients with stage III NSCLC also showed a lower physical activity than healthy subjects (approximately \(6 \text{~v.~} 20\)\(^{50}\)), and patients with pre-cachexia and cachexia showed a non-significant lower physical activity than no-cachexia patients.

When using the selected frameworks, we encountered some issues. First, patients with weight loss as well as complaints and/or inflammation were incorrectly justified as having no cachexia by the general framework, which requires three positive scores on complaints and inflammation. Patients with \(\geq 5\%\) weight loss, in combination with two positive scores on complaints and inflammation, were not classified as cachectic. Also, the ESPEN Special Interest Group did not classify these patients as pre-cachectic, as their weight loss was more than \(5\%\). Second, cut-off points for anorexia, CRP and FFM index were lacking for the pre-cachexia and cancer-specific frameworks. For pre-cachexia, weight loss \(\leq 5\%\) was described, but it was unclear if this accounted for patients with a weight loss of \(0\%\). We solved these issues by consulting the authors. In line with current knowledge, we found a positive correlation between pro-inflammatory indexes (CRP and serum IL-6), and these were negatively correlated with Hb and serum albumin. Interestingly, when other cut-off points for inflammatory parameters were applied, we observed the same presence of (pre)cachexia.

Validation of cachexia instruments in large groups of patients with cancer is still required, but a ‘gold standard’ is lacking. The association of cachexia with survival is informative, but validation of instruments against one or more indicators of cachexia (e.g. standardised assessment of muscle mass) is preferable. Further studies in larger populations are warranted to validate these new instruments and to more extensively explore the prognostic value in patients with cancer. Ideally, worldwide cancer centres record a number of biomarkers and cachexia parameters, follow-up treatment adherence and survival, and merge these data in order to validate definitions and their prognostic value. A promising parameter might be proteolysis-inducing factor, which has been found in the urine of cachectic patients with cancer\(^{49}\).

In conclusion, new consensus-based frameworks show that pre-cachexia and cachexia are prevalent in patients with stage III NSCLC. Cachexia appears to be associated with a shorter overall survival and a reduced quality of life.

**Supplementary material**

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**References**


