Nimotuzumab combined with radiotherapy for esophageal cancer: preliminary study of a Phase II clinical trial

Objective: To determine the safety and therapeutic effects of nimotuzumab (h-R3) combined with radiotherapy in esophageal cancer.

Methods: This Phase II clinical trial involved 42 patients with stage II (inoperable or refused surgery) to stage IV (supraclavicular lymph node metastasis only) esophageal cancers treated between November 2008 and July 2010. All patients had squamous cell carcinomas, and all received three-dimensional conformal radiotherapy and 200 mg nimotuzumab per week during radiotherapy.

Results: There were 9, 25, and 8 patients with stage II, III and IV disease, respectively. All except two patients received 50–70 Gy radiation; 37 patients (88.1%) received more than five nimotuzumab doses. Grade III toxicities (21.4% of all adverse events) included esophagitis and gastrointestinal, dermatological and hematological toxicities. Complete response, partial response, stable disease, and progressive disease were observed in 0, 22 (52.4%), 17 (40.5%) and 3 (7.1%) patients at 1 month after the treatment. The epidermal growth factor receptor (EGFR) overexpression rate was 95.2%. After a median follow-up of 37 months, the median survival time (MST) was 5 months. The 2 year and 3 year overall survival (OS) rates were 33.3% and 26.2%, respectively. The median progression-free survival (PFS) time was 10 months. The 2 year and 3 year PFS rates were 24.5% and 22.1%, respectively. The MST in the 15 patients with (+++) EGFR expression (group A) and 7 patients with (++) EGFR expression (group B) was 15 and 11 months, respectively. The 2 year and 3 year OS rates were 46.2% and 38.5% in group A and 28.6% and 28.6% in group B, respectively (P = 0.405).

Conclusion: Although concurrent chemoradiotherapy was the standard care for locally advanced esophageal cancer, radiotherapy was the choice for those who were refused or could not tolerate chemoradiotherapy. Our study shows that nimotuzumab combined with radiotherapy was well tolerated in patients with esophageal cancer. EGFR overexpression was more common than previously reported. OS was higher after combined therapy than after historical control radiotherapy alone. Further studies are required to confirm the therapeutic efficacy of nimotuzumab in esophageal cancer.

Keywords: esophageal neoplasms, nimotuzumab, radiotherapy, targeted therapy, treatment outcomes

Introduction

Esophageal cancer is the sixth leading cause of cancer-related deaths in the world.1,2 The morbidity and mortality rates of esophageal cancer in the People’s Republic of China are the highest in the world, and over 50% of patients have locally advanced or metastatic disease at presentation.3 Currently, patients who are unfit to undergo surgery or have locally advanced esophageal cancer are offered concurrent chemoradiotherapy.4,6
Modern approaches to cancer treatment are focused on combination strategies involving surgery, chemotherapy, radiotherapy, new chemotherapy drugs and targeted therapy.

Many malignant tumors overexpress epidermal growth factor receptor (EGFR), which initiates signal transduction by activating a receptor-associated tyrosine kinase. EGFR overexpression has been associated with cancer cell invasion, metastasis, and poor prognosis. EGFR inhibitors have proven efficacious in patients with non-small cell lung, colon, intestinal, and head and neck cancers.

Nimotuzumab (h-R3) is a humanized monoclonal antibody. In preclinical studies, nimotuzumab has shown significant antitumor, proapoptotic and antiangiogenic activities. In an in vitro study, nimotuzumab was shown to radiosensitize a non-small cell lung cancer cell line. In another, more recent study, nimotuzumab enhanced the effects of radiotherapy on esophageal squamous cell carcinoma cells with a functional and active EGFR pathway. Several Phase I and II clinical studies have confirmed the safety profile of the combination of nimotuzumab with radiotherapy. Nimotuzumab combined with radiotherapy or chemoradiotherapy has proven clinically beneficial, improving overall survival (OS) and progression-free survival (PFS) in patients with head and neck cancer.

Between November 2008 and July 2010, we conducted a Phase II clinical trial in which patients with stage II (inoperable or refused surgery) to stage IV (supraclavicular lymph node metastasis only) esophageal cancer received nimotuzumab in combination with radiotherapy. Most of the patients in our study had previously refused chemotherapy, and others could not tolerate chemotherapy. The purpose of this multicenter, Phase II study was to determine the safety and therapeutic effect of nimotuzumab in combination with radiotherapy in patients with esophageal cancer.

The trial was approved by the China State Food and Drug Administration (CFDA). The study protocol was approved by the ethical review committee, and permission to conduct the study was granted by the institutional review board. The trial has been registered according to Good Clinical Practice at the Cancer Institute and Hospital, Chinese Academy of Medical Sciences (CAMS). All enrolled patients in the trial voluntarily signed the informed consent forms.

**Methods**

**Endpoints**

The endpoints were toxicity, clinical response, PFS, and OS.

**Patient selection and enrollment**

In total, 120 esophageal cancer patients from all across the People’s Republic of China were screened. Of these, 42 patients with histologically confirmed stage II (inoperable or refused surgery) to stage IV (supraclavicular lymph node metastasis only) cancer were eligible and were assigned to this trial, which was conducted in six hospitals (Cancer Hospital, CAMS; Cancer Hospital, Harbin Medical University; Tongji Cancer Center Hospital; Cancer Hospital, Tianjin Medical University; Liao-Ning Province Cancer Hospital; Beijing Hospital) between November 2008 and July 2010.

The inclusion criteria were as follows: age, 18–75 years; thoracic-segment esophageal cancer in stages II (inoperable or refused surgery) to IV (supraclavicular lymph node metastasis only); no history of surgery, although history of radio- or chemotherapy was acceptable; estimated survival time, ≥3 months; Karnofsky performance score, >60; no serious disease of major organs that would affect physical function; and voluntary signing of consent forms.

The exclusion criteria were as follows: pregnancy or lactation, history of other malignant disease, and joining another clinical trial prior to the study. Of the 42 eligible patients, 34 were male and 8 were female; their median age was 63 years (range, 35–74 years). Table 1 shows further patient characteristics.

All eligible patients underwent endoscopy, biopsy, and computed tomography (CT) of the neck, chest, and abdomen, esophagography, and other physical examinations, including blood tests. Patients with metastases, involvement of more than one lymph node, and lymph nodes ≥0.5 cm in diameter were eligible. Radionuclide bone scans and brain magnetic resonance imaging were offered to the patients. Other tests, such as endoesophageal ultrasonography and positron emission tomography–computed tomography (PET-CT) were widely available at the time of the trial. EGFR expression was recommended to be measured in all samples, if possible.

**Treatment schedule**

**Radiotherapy**

All patients received three-dimensional conformal radiotherapy with 4–10 Mv X-ray accelerators. A CT simulation with a slice thickness of 5 mm was obtained before irradiation. The gross tumor target volume (GTV) was defined as the primary tumor, and the GTVnd as the enlarged regional lymph nodes. The clinical target volume (CTV) was defined as the GTV plus an area bounded by a margin of 3 cm in the superior and inferior directions to the GTV, 6 mm in the left
and right directions and 6 mm in the anterior and posterior directions. The CTV also included the GTVnd and the area of the prophylactic regional lymph node. In the case of upper-segment and middle-segment thoracic esophageal cancers, the superior margin of the prophylactic regional lymph node was the upper border of the T1 vertebra, and the inferior margin was the subcarinal region. In the case of lower-segment cancers, the superior margin of the prophylactic regional lymph node was the thoracic inlet, and the inferior margin was the left gastric region. The planning target volume (PTV) was defined as the CTV plus an area bounded by a margin of 5 mm extending three-dimensionally around the CTV. The total dose to 95% of the PTV was 50–60 Gy, which was administered in 2-Gy once-daily fractions for 5 days a week over 5–6 weeks.

**Targeted therapy**

In a clinical trial of the humanized anti-epidermal growth factor receptor monoclonal antibody h-R3 with concurrent chemoradiotherapy for unresectable head and neck carcinomas reported by Crombet et al, it was found that a 200 mg/week of h-R3 (the lab code for nimotuzumab) was the appropriate dose on the basis of h-R3 serum levels and the response and survival. Therefore, all patients in our trial received 5–6 once-weekly injections of 200 mg h-R3 diluted in 250 mL 0.9% sodium chloride, administered as an intravenous infusion over 1 hour.

**Follow-up**

All patients were monitored weekly during the treatment for signs of acute toxicity. Follow-up was conducted at 3 month intervals in the first 3 years and at 6 month intervals thereafter.

**Evaluation of therapeutic effect and toxicity**

The endpoints for the trial were toxicity, clinical response, PFS, and OS. Toxic effects were assessed once a week, using the National Cancer Institute’s Common Toxicity Criteria, version 3.0. Clinical response was assessed using the Response evaluation criteria in solid tumors (RECIST). Two-dimensionally measurable disease was defined by the length and width of the primary tumor and the diameter of the largest positive lymph node. Response in assessable disease was determined by at least two observers. OS was defined as the interval between the date of treatment commencement and the date of death from esophageal cancer or treatment-related causes. Patients who died from other causes were excluded from the analysis. The rates of adverse events and toxic effects, and the OS were determined. All case report forms were assessed centrally in the Cancer Hospital, CAMS.

**Statistical analysis**

We performed an intention-to-treat analysis. SPSS software, version 11.5 (IBM Corporation, Armonk, NY, USA), was used. Analyses were also conducted for subgroups based on age, gender, performance status, tumor location, pathological type, EGFR expression, and tumor length and width. Survival time was calculated from the date of the first treatment until the date of death. PFS and OS data were estimated with the Kaplan-Meier method, and EGFR expression subgroups were compared using the log-rank test.

**Results**

**Treatment outcomes**

**Toxicity**

The common acute toxicities during the treatment were esophagitis, and blood/bone marrow, dermatological, and gastrointestinal complications; Table 2 shows the details by grade and incidence. Other toxicities included grade 1 paresthesia (1/42), septicemia (1/42), herpes virus infection (1/42), hyperbilirubinemia (1/42), pulmonary infection (3/42),
Table 2 Common acute toxicities during treatment

<table>
<thead>
<tr>
<th></th>
<th>Grade 1</th>
<th>Grade 2</th>
<th>Grade 3</th>
<th>Grade 4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Esophagitis</td>
<td>16 (38.1%)</td>
<td>20 (47.6%)</td>
<td>3 (7.1%)</td>
<td>0</td>
</tr>
<tr>
<td>Blood/bone marrow</td>
<td>14 (33.3%)</td>
<td>10 (23.8%)</td>
<td>1 (2.4%)</td>
<td>0</td>
</tr>
<tr>
<td>Dermatological</td>
<td>15 (35.7%)</td>
<td>4 (9.5%)</td>
<td>4 (9.5%)</td>
<td>0</td>
</tr>
<tr>
<td>Gastrointestinal</td>
<td>6 (14.3%)</td>
<td>2 (4.8%)</td>
<td>1 (2.4%)</td>
<td>0</td>
</tr>
</tbody>
</table>

severe fatigue (2/42), fever (9/42) and tracheoesophageal fistula (1/42). Only one patient was allergic to nimotuzumab.

The incidence of grade 3 toxicity was 21.4%; no grade 4 toxicity occurred. Mild, nimotuzumab-related skin rash was observed in four patients but did not require treatment.

All patients received 50–60 Gy of irradiation, except for two patients who received only 46 Gy (one patient refused to continue treatment, and the other patient requested surgery), and one patient who received 70 Gy. In all, 37 (88.1%) patients received five or six doses of 200-mg nimotuzumab, and five patients received this dose fewer than 5 times because of adverse effects: allergy, pulmonary infection, severe fatigue, or tracheoesophageal fistula.

Therapeutic effects

The date of the last follow-up was January 30, 2013. The median follow-up time was 37 months (range 31–41 months). At 1 month after the treatment, none of the patients had attained complete response (CR); 22 (52.4%) patients had achieved partial response (PR); whereas 17 (40.5%) and 3 (7.1%) patients showed stable disease (SD) and progressive disease (PD), respectively. The objective clinical response was evaluated at the last follow-up, and CR, PR, SD, and PD were observed in 7 (16.7%), 3 (7.1%), 0 (0%), and 32 (76.2%) patients, respectively. Six patients had achieved CR by the sixth month after treatment, and another patient attained CR by the seventh month. The median survival time (MST) was 14 months, and the median PFS was 10 months. The 2 year and 3 year OS rates were 33.3% and 26.2%, respectively, and the corresponding PFS rates were 24.5% and 22.1% (Figure 1).

Univariate and multivariate analyses showed that gender, age ≥65 years, stage IV disease, lower-segment tumors, and tumor width and length were not predictors of OS and PFS. Local recurrence and distant metastases were observed in 20 (47.6%) and 14 (33.3%) patients, respectively. Twelve metastases were detected in the lungs, three in the bones, three in the liver, two in the pleura, one in the hypodermis, and one in a distant lymph node. At the last follow-up, 31 (73.8%) patients had died: 16 (38.1%) of local recurrence, 13 (31%) of metastases, and 2 (4.8%) of other causes. The median disease-specific survival time was 14 months. The 2 year and 3 year disease-specific survival rates were 32.7% and 32.7%, respectively.

EGFR expression

EGFR expression was determined in 21 patients; biopsy specimens were obtained from these patients via esophagoscopy and subjected to an immunochemical assay. The immunoreactivity of EGFR was graded into four groups according to the intensity of cell membrane EGFR staining in the whole tumor: high (+++), markedly stronger staining than normal esophageal epithelium; medium (++), moderately stronger staining than normal esophageal epithelium; low (+), staining identical to that of normal epithelium (Figure 2); and negative (−), faint staining. Strong (+++) and moderate (+++) staining indicated EGFR overexpression, and were found in 20 patients (95.2%). EGFR expression at levels (+++), (+++) and (+) was found in 13, 7, and 1 patient, respectively. The MST in the 13 patients with (+++) EGFR expression (group A) and the 7 patients with (+++) EGFR expression (group B) was 15 and 11 months, respectively. The 2 year and 3 year OS rates were 46.2% and 38.5%, respectively, in group A and 28.6% and 28.6%, respectively, in group B; the between-group differences were not significant (P = 0.405; Figure 3).

Discussion

In our study of esophageal cancer patients, nimotuzumab combined with radiotherapy was well tolerated. The incidence of grade 3 toxicity was 21.4%; no grade 4 toxicity occurred. Mild nimotuzumab-related rash occurred, but nimotuzumab-related diarrhea did not occur. Only one patient was allergic to nimotuzumab. Nimotuzumab did not seem to increase the acute toxicity of radiotherapy. Safran et al have reported incidence rates of 23%, 15%, and 5% for skin...
rash, esophagitis and hypersensitivity of grades 3 or higher, respectively, in patients receiving HER2-targeted therapy for esophageal tumors.21

In our study, most patients who achieved CR did so at 6 months after the treatment, suggesting a delayed therapeutic effect. This finding might indicate that the best time to evaluate the objective clinical response, especially CR, is at 6 months after treatment.

The 2 year and 3 year OS rates were 33.3% and 26.2%, respectively, and the corresponding PFS rates were 24.5% and 22.1%. The MST was 14 months. These results are more favorable than those reported for radiotherapy alone. In a study comparing concurrent chemoradiotherapy and radiotherapy alone in esophageal cancer patients with lymph node metastasis, the 2 year and 3 year OS rates in the radiotherapy-alone group were 38.5% and 18.5%, respectively.22 In the Radiation Therapy Oncology Group 85-01 trial, no patient was alive in the radiotherapy-alone group after 3 years; the MST in this group was 8.9 months.2 In an Eastern Cooperative Oncology Group trial, the MST was 9.2 months and the 3 year OS was 8% in the radiotherapy-alone group.23

For patients with locally advanced esophageal cancer (those who are inoperable or who refuse surgery), concurrent chemoradiotherapy is the standard of care. Most patients in our study refused chemotherapy, and others could not tolerate chemotherapy. For such patients, radiotherapy alone seems to be the only treatment currently accepted/accepted. However, our study showed that combined radiotherapy and targeted therapy could improve the OS over that achievable with radiotherapy alone.

The rate of EGFR overexpression was 95.2% in our study, and this was higher than the rates previously reported. EGFR overexpression has been observed in 29%-90% of esophageal cancers and may be correlated with poor prognosis and response.24 EGFR overexpression has also been associated with invasion, metastasis and poor prognosis,8 and with chemo- and radio-resistance.25-27 In our study, group A patients (EGFR, ++++) had a higher OS and MST than group B patients (EGFR, ++). Although the differences between the two groups were insignificant, this finding might be attributable to the small number of patients. Moreover, this result may indicate that the effectiveness of EGFR
Table 3 Summary of trials of targeted therapy for esophageal cancer

<table>
<thead>
<tr>
<th>Year</th>
<th>Authors</th>
<th>Path</th>
<th>NP</th>
<th>NEP</th>
<th>TS</th>
<th>RR*</th>
<th>M/O</th>
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<td>AC</td>
<td>17</td>
<td>5</td>
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<td>Trastuzumab + Cis/Pacl/RT</td>
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<td>SCC</td>
<td>10</td>
<td>13</td>
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<td>17</td>
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<td>AC</td>
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<td>37</td>
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<td>PCR</td>
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<tr>
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<td>SCC</td>
<td>8</td>
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<td>AC</td>
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<td>45</td>
<td>Cetuximab + Carb/Pacl/RT</td>
<td>27</td>
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<tr>
<td>2008</td>
<td>De Vita et al</td>
<td>SCC</td>
<td>9</td>
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<td>18</td>
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<td>72.2</td>
<td>2y 44.4</td>
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</table>

Abbreviations: AC, adenocarcinoma; SCC, squamous cell carcinoma; Carb, carboplatin; Pacl, paclitaxel; rT, radiotherapy; FOLFOX, weekly 5-Fu/oxaliplatin/leucovorin infusion; 5-Fu, 5-fluorouracil; Cis, cisplatin; Doc, docetaxel; CPT, irinotecan; Path, pathology; NP, number of patients; NEP, number of esophageal cancer patients; TS, treatment strategy; RR*, response rate (%); M/O, median survival time (years/months)/overall survival (%).

The treatment strategies were also different and included targeted therapy alone; combination chemotherapy and targeted therapy; combination radiotherapy and targeted therapy; and combination concurrent chemoradiotherapy and targeted therapy. The drugs used for targeted therapy also differed among these trials. The number of patients in these trials was small. Although some trials showed encouraging results, targeted therapy for esophageal cancer requires further evaluation.

Conclusion

Although concurrent chemoradiotherapy has been the standard care for locally advanced esophageal cancer, radiotherapy was the choice for those who were refused or could not tolerate chemoradiotherapy. Our study shows that patients with esophageal cancer tolerated nimotuzumab combined with radiotherapy well. The OS after combined h-R3 treatment and radiotherapy was higher than that after historical control radiotherapy alone. EGFR overexpression was common. Further confirmatory studies are required.
Acknowledgment

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Disclosure

The authors of this study declare no conflicts of interest. The sponsor of this study had no role in the study design, data collection, data analysis or data interpretation.

References


Nimotuzumab plus RT for esophageal cancer

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