Efficacy and safety of amrubicin hydrochloride for treatment of relapsed small cell lung cancer

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Abstract: Long-term survival is quite uncommon in refractory small cell lung cancer (SCLC) patients, with less than 25% of patients with limited-stage disease and 1%–2% of patients with extensive-stage disease remaining alive at five years. Recent clinical studies have demonstrated the promising efficacy of amrubicin for patients with relapsed SCLC. This review presents the results of clinical studies showing the efficacy and safety of amrubicin for the treatment of relapsed SCLC. Amrubicin is a synthetic anthracycline agent with a similar structure to doxorubicin, in which the hydroxyl group at position 9 in amrubicin is replaced by an amino group to enhance efficacy. It is converted to an active metabolite, amrubicinol, which is 5–54 times more active than amrubicin. Amrubicin and amrubicinol are inhibitors of DNA topoisomerase II, exerting their cytotoxic effects by stabilizing a topoisomerase II-mediated cleavable complex. The toxicity of amrubicin is similar to that of doxorubicin, although amrubicin shows almost no cardiotoxicity. In the relevant trials, amrubicin was administered intravenously at a dose of 35–40 mg/m² on days 1–3 every three weeks. The response rate was 34%–52% and median survival times were 8.1–12.0 months. Common hematologic toxicities included neutropenia, leucopenia, anemia, thrombocytopenia, and febrile neutropenia. Nonhematologic adverse events included Grade 3–4 anorexia, asthenia, hyponatremia, and nausea. The results of the studies which demonstrated the efficacy of monotherapy for relapsed SCLC involved mainly Japanese patients. Therefore, it is necessary to conduct more clinical studies in non-Japanese patients to confirm the efficacy of amrubicin.

Keywords: amrubicin, amrubicinol, small cell lung cancer, relapse

Introduction

Small cell lung cancer (SCLC) accounts for approximately 15% of all lung cancer cases, with two-thirds of patients presenting with extensive disease (ED). Without treatment, tumor progression in patients with SCLC is rapid, with a poor prognosis. However, the disease shows a high response rate to chemotherapy and radiotherapy, except in a low percentage of patients. Treatment options for relapsed SCLC patients remain limited. A randomized trial demonstrated that single-agent topotecan was at least as efficacious as the three-drug combination of cyclophosphamide-doxorubicin-vincristine for the treatment of patients with sensitive relapsed cases.1 Response rates and median survival times were 24% and 25.0 weeks for topotecan, and 18% and 24.7 weeks for cyclophosphamide-doxorubicin-vincristine, respectively. In previously untreated ED-SCLC, amrubicin yielded an extremely high response rate of 79% and a median survival time of 11 months, which was comparable with the results achieved with
platinum.Recently, clinical studies have demonstrated the efficacy of amrubicin in patients with relapsed SCLC. This review presents the results of clinical studies showing the efficacy and safety of amrubicin for the treatment of relapsed SCLC.

**Structure and characteristics**

Amrubicin hydrochloride is a synthetic anthracycline agent, with a similar structure to doxorubicin, in which the hydroxyl group at position 9 group in amrubicin is replaced by an amino group to enhance efficacy (see Figure 1). Amrubicin is converted to an active metabolite, amrubicinol, which is 5–54 times more active than amrubicin, through reduction of its C-13 ketone group to a hydroxyl group by carbonyl reductase. Other enzymes metabolizing amrubicin and amrubicinol are nicotinamide adenine dinucleotide phosphate (NADPH)-dependent P450 reductase and NAD[P]H-dependent quinone oxidoreductase. Doxorubicin, amrubicin, and amrubicinol are inhibitors of DNA topoisomerase II, exerting their cytotoxic effects by stabilizing a topoisomerase II-mediated cleavable complex. In addition, they are more or less only one-tenth as potent as doxorubicin in producing DNA intercalation.

**Preclinical studies**

Antitumor activity and toxicologic aspects were first reported for amrubicin by Morisada et al. They evaluated this agent in six murine systems and nine human tumor-nude mouse systems, and found that the antitumor activity of amrubicin was superior to adriamycin in human tumor xenografts, and almost equal against murine experimental tumors. They also evaluated toxicity in mice after a bolus intravenous injection. The acute toxic signs were body weight decrease, ataxia, hair loss, and myelosuppression, and these toxicities were qualitatively comparable with those induced by adriamycin. The maximum tolerated dose was estimated to be 25 mg/kg in four mouse strains, and the drug had anticancer activity against human lung cancer xenografts *in vivo.*

Cardiomyopathy is a burdensome toxicity with the anthracyclines. Suzuki et al evaluated the degree of cardiotoxicity of amrubicin compared with that of adriamycin in rabbits. The drugs were intravenously administered three times a week for eight weeks. In this study, prolongation of the QT₅₀ interval and STT changes were observed in rabbits administered amrubicin and adriamycin. Morphologic studies showed that myocardial tissue damage in animals administered amrubicin was comparable with that seen in controls. Considering the results of the antitumor efficacy studies comparing amrubicin with adriamycin, they concluded that the cardiotoxicity of amrubicin was very slight.

**Relapsed small cell lung cancer**

Long-term survival is quite uncommon in refractory SCLC patients, with less than 25% of patients with limited-stage disease and 1%–2% of patients with extensive-stage disease remaining alive at five years. A Phase II study was conducted in patients with relapsed disease who had previously received one or two regimens, including at least one regimen of platinum-based chemotherapy (Table 1). Sixty patients were enrolled in this multicenter trial. The disease progressed within 60 days after the final dose of previous chemotherapy in 16 and 44 refractory patients, respectively. The sensitive groups, in which complete response (CR) or partial response (PR) was observed with previous chemotherapy.
Amrubicin in relapsed lung cancer

Amrubicin was administered intravenously at a dose of 40 mg/m² on days 1–3 every three weeks. The response rate was 52% (95% confidence interval [CI] 38%–65%). There were no differences in the response rate, ie, 50% (95% CI 25%–75%) for refractory disease and 52% (95% CI 37%–68%) for sensitive disease. The median survival times were 10.3 months in the refractory group and 11.6 months in the sensitive group, respectively (P = 0.0974, log rank test). Common adverse events were hematologic toxicities, including Grade 3–4 neutropenia (83%), leucopenia (70%), anemia (33%), thrombocytopenia (20%), and febrile neutropenia (5%). Nonhematologic adverse events included Grade 3–4 anorexia (15%), asthenia (15%), hyponatremia (8%), and nausea (5%).

Another Phase II study of amrubicin in patients with previously treated SCLC was conducted by Kaira et al10 (Table 2). Twenty-nine patients with relapsed SCLC who had previously received platinum-based chemotherapy were enrolled in the trial, including 10 patients with sensitive relapse and 19 patients with refractory relapse. Amrubicin was administered intravenously at a dose of 35 mg/m² on days 1–3 every three weeks. The response rate was 44.8% (95% CI 26%–64%), being 60% for sensitive cases and 37% for refractory cases. No significant difference in the response rate was observed between sensitive cases and relapsed cases (P = 0.233, log rank test). The median progression-free survival and median survival times were 4.0 months (sensitive relapse, 4.0 months; refractory relapse, 4.0 months) and 12.0 months (sensitive relapse, 12.0 months; refractory relapse, 11.0 months), respectively. There was no difference in median progression-free survival and median survival time between sensitive relapse and refractory relapse. Grade 3 or 4 neutropenia and febrile neutropenia were observed in 42% and 3% of patients, respectively. Nonhematologic toxicity higher than Grade 3 was not observed. The results of this study show the efficacy of monotherapy for relapsed SCLC. However, this study involved only Japanese patients, so it would be necessary to conduct clinical studies in non-Japanese patients to confirm efficacy.

A randomized Phase II trial of amrubicin versus topotecan as second-line treatment for sensitive ED-SCLC was therefore conducted11 (Table 3). Seventy-six patients who had previously received platinum-based frontline chemotherapy were enrolled. All were sensitive cases, in which CR or PR had been observed with previous chemotherapy and the disease had then progressed or relapsed within at least 90 days of the final dose. Patients were randomized at a 2:1 ratio to receive either amrubicin or topotecan. Amrubicin was administered intravenously at a dose of 40 mg/m² on days 1–3 every three weeks. Topotecan was administered intravenously at a dose of 1.5 mg/m² on days 1–5 every three weeks. The response rate for amrubicin was 34% (95% CI 22%–48%), and for topotecan was 4% (95% CI 1%–19%). There was a trend towards a longer

### Table 1 Phase II trial of amrubicin for treatment of refractory or relapsed small cell lung cancer (Thoracic Oncology Research Group Study 0301)1

<table>
<thead>
<tr>
<th>Patients (n)</th>
<th>Sensitive cases</th>
<th>Refractory cases</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>CR</td>
<td>1</td>
<td>1</td>
<td>2</td>
</tr>
<tr>
<td>PR</td>
<td>22</td>
<td>7</td>
<td>29</td>
</tr>
<tr>
<td>SD</td>
<td>10</td>
<td>2</td>
<td>12</td>
</tr>
<tr>
<td>PD</td>
<td>11</td>
<td>6</td>
<td>17</td>
</tr>
<tr>
<td>Response rate (% 95% CI)</td>
<td>52 (37–68)</td>
<td>50 (25%–75%)</td>
<td>52 (38%–65%)</td>
</tr>
<tr>
<td>PFS (months, 95% CI)</td>
<td>42 (10.0–15.8)</td>
<td>2.9 (1.4–4.6)</td>
<td>3.9 (3.4–4.6)</td>
</tr>
<tr>
<td>MST (months, 95% CI)</td>
<td>11.6 (10.0–15.8)</td>
<td>10.3 (4.8–∞)</td>
<td>11.0 (10.0–13.2)</td>
</tr>
<tr>
<td>One-year survival (% 95% CI)</td>
<td>45.5 (29.9–59.8)</td>
<td>40.3 (15.1–64.6)</td>
<td>44.1 (30.6–56.8)</td>
</tr>
</tbody>
</table>

**Abbreviations:** CI, confidence interval; CR, complete response; PR, partial response; SD, stable disease; PD, progressive disease; MST, median survival time; PFS, progression-free survival.

### Table 2 Phase II trial of amrubicin for treatment of relapsed small cell lung cancer10

<table>
<thead>
<tr>
<th>Patients (n)</th>
<th>Sensitive cases</th>
<th>Refractory cases</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Response rate (% 95% CI)</td>
<td>60 (26–64)</td>
<td>37 (26–64)</td>
<td>45 (26–64)</td>
</tr>
<tr>
<td>Progression-free survival (months)</td>
<td>4</td>
<td>4</td>
<td>4</td>
</tr>
<tr>
<td>Median survival time (months)</td>
<td>12</td>
<td>11</td>
<td>12</td>
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</tbody>
</table>
amrubicin was 17% (95% CI 2%–48%), and for topotecan was 0% (95% CI 0%–28%). The median progression-free survival time with amrubicin was 3.5 months and with topotecan was 2.2 months. The median overall survival time with amrubicin was 8.1 months and with topotecan was 8.4 months. There was no difference in the frequency of hematologic toxicity more than Grade 3 between amrubicin and topotecan. These studies show that amrubicin monotherapy is an encouraging regimen for second-line treatment of SCLC.

**Conclusion**

Clinical investigation of the novel anticancer agent, amrubicin, has increased quickly, and there are high expectations for this agent in trials to improve the outcome for relapsed SCLC patients. Amrubicin is an active agent for the treatment of relapsed SCLC, but because it is strongly myelotoxic, particular care should be taken with its use.

**Disclosure**

The authors declare no potential conflicts of interest.

**References**


