Meta-analytic evaluation of the correlation between event-free survival and overall survival in randomized controlled trials of newly diagnosed Ewing sarcoma.

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Abstract

Background

In randomized controlled trials (RCTs) of adjuvant treatment for malignant tumors, event-free survival (EFS) is considered the most acceptable surrogate for overall survival (OS). However, even though EFS has repeatedly been selected as a primary endpoint in RCTs of Ewing sarcoma (ES), the surrogacy of EFS for OS has not been investigated. This study aimed to evaluate the surrogacy of EFS for OS in RCTs of chemotherapy for newly diagnosed ES using a meta-analytic approach.

Methods

We identified seven RCTs of newly diagnosed ES through a systematic review, and a meta-analysis was performed to evaluate the efficacy and adverse events associated with chemotherapy for previously untreated ES. The correlation between EFS and OS was investigated using weighted linear regression analysis and Spearman rank correlation coefficients ($\rho$). The strength of the correlation was evaluated using the coefficient of determination ($R^2$).

Results

A total of 3,612 patients were randomly assigned to 17 treatment arms in the eligible RCTs. The meta-analysis revealed that the hazard ratios for OS and EFS showed significantly better results in the experimental treatment groups with increasing toxicities. The correlation between the hazard ratios for EFS and OS was good ($R^2 = 0.747$, $\rho = 0.683$), and the correlation tended to be more favorable in cases of localized ES ($R^2 = 0.818$, $\rho = 0.929$).

Conclusions

Overall, the trial-level correlation between EFS and OS was good for newly diagnosed ES and was very good in cases of localized disease. EFS may be useful as a surrogate endpoint in RCTs of ES, and the surrogacy of EFS is worth verifying using individual data.

Background

Ewing sarcoma (ES) is the second most frequent malignant bone tumor in children. However, it accounts for only 1% of all childhood cancers in the United States [1]. ES is a highly malignant disease, involving the dense proliferation of round tumor cells without specific histologic
differentiation. Tumors presenting a similar histological appearance also occur in soft tissues. However, such tumors are considered to be the same as ES arising in the bone because they present the same chromosomal abnormality and fusion gene [1]. Intensive multidrug chemotherapy and definitive local control are the standard treatment options for ES [2].

A standard regimen for newly diagnosed ES has been examined in some randomized controlled trials (RCTs). RCTs provide pivotal evidence for determining standard treatments, and primary endpoints in RCTs are considered the most important measures for judging the effectiveness of treatment options; this influences trial conclusion, and subsequently, patient survival. Overall survival (OS) is the preferred primary endpoint in RCTs, as it is an index that eliminates subjectivity and ambiguity because it can be clearly defined and judged. Additionally, because improvement in OS is the primary objective of cancer treatment, OS is an appropriate indicator for evaluating treatment efficacy. However, using OS as the primary endpoint in RCTs entails longer durations, larger sample sizes, and higher costs. In addition, multiple lines of new drugs may extend patients’ post-progression survival, making it difficult to assess the true effects of the regimens used in the trial and the influence of post-protocol treatment. Event-free survival (EFS) has, therefore, been used as the primary endpoint in many RCTs of ES. However, the correlation between EFS and OS in RCTs of ES has not yet been investigated.

Accordingly, in this study, data on all RCTs that involved chemotherapy for newly diagnosed ES were collected, and the correlation between EFS and OS was evaluated by using a meta-analysis.

**Methods**

**Study selection and data extraction**

Based on the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines [3], we conducted a systematic search of PubMed, Scopus, EBSCOhost MEDLINE, and the Cochrane Central Register of Controlled Trials. We searched for all RCTs of ES published in English between January 1973 and October 2018. The study inclusion criterion was phase II or III RCTs of systemic chemotherapy for newly diagnosed ES without prior treatment. We excluded non-randomized clinical trials, reviews, and meta-analyses. RCTs retrieved by this search were screened independently by two
authors and cross-checked (KT and MK).

Data extraction of the date of publication, trial name, patient accrual period, phase of study, primary and secondary endpoints, regimens and doses used in the standard and experimental arms, number of patients, sex, age, number of metastatic cases, description of intention-to-treat (ITT) analysis, description of post-protocol treatment, radiological and histological responses to chemotherapy, survival data, and adverse events (AEs) was performed.

The medians, hazard ratios (HRs), 95% confidence intervals (CIs), and \( P \)-values were extracted for OS and EFS. The pathological response rate was defined as the proportion of assessed patients showing >90% tumor necrosis in the resected specimens. Data on 1-, 3-, and 5-year EFS and 1-, 3-, and 5-year OS were extracted based on Kaplan-Meier estimates. If these data were not described, Kaplan-Meier curves of EFS or OS were used for the estimation as binary proportions. Data were extracted and cross-checked by two authors (KT and MK). In cases of discrepancies between these two authors, other authors (TI or II) were consulted to reach a consensus.

**Statistical analysis**

Pooled HRs and their corresponding 95% CIs were subjected to meta-analyses, and the values for EFS and OS were obtained. The odds ratios (ORs) and corresponding 95% CIs for 1-, 3-, and 5-year EFS and 1-, 3-, and 5-year OS were also calculated. Meta-analyses were performed using inverse-variance and a Mantel-Haenszel random- or fixed-effect model. The random-effect model was used when the \( p \)-value in the heterogeneity test was <0.1. Heterogeneity was evaluated by Cochrane’s Q-test and \( I^2 \) statistics. Meta-analyses were conducted using Review Manager software (version 5.3; Nordic Cochrane Centre, Cochrane Collaboration, Copenhagen, Denmark).

The association between EFS and OS was evaluated using a weighted linear regression test with the study sample size. The correlation between HRs for the surrogate endpoints and OS was assessed using Spearman’s rank correlation coefficients (\( \rho \)). The strength of the association was also investigated using the coefficient of determination (\( R^2 \)) [4,5]. Coefficient values >0.9 were defined as excellent, >0.75 as very good, >0.5 as good, >0.25 as moderate, and \( \leq 0.25 \) as poor [6]. Sensitivity
analyses were performed to eliminate the treatment arms of high-risk and metastatic disease from
the evaluation of surrogacy. Further sensitivity analyses were conducted by eliminating two old RCTs
(IESS-I and -II).
Other statistical analyses were conducted using SAS (version 9.4; SAS Institute, Cary, NC, USA). \( P \)-
values reflected two-sided tests, with \( P < 0.05 \) indicating statistical significance.

Results

**Characteristics of RCTs included in the analysis**

In this systematic literature search, 2,432 articles were identified. After excluding 50 duplicates,
2,382 studies were further screened. The full texts of 28 articles were finally evaluated after
excluding 2,354 studies. Of the 28 studies, 3 duplicate publications, 6 repeat publications, 10 non-RCT
studies, and 2 non-chemotherapeutic studies were excluded. The remaining 7 RCTs were considered
eligible for the meta-analysis (Figure 1). The characteristics of the eligible RCTs are summarized in
Additional files 1: Table S1 [7-13]. In the 7 eligible RCTs, 3,612 patients were randomly assigned to 17
treatment arms. All treatment arms comprised combination regimens with 3–5 cytotoxic drugs. One
treatment arm included lung irradiation as the protocol treatment. No study included molecular-
targeted therapy or immune therapy. Study phases and post-protocol treatments were not clearly
described in any study, and ITT analyses were conducted only in 2. The primary endpoint was defined
in 5 of 7 RCTs as EFS, including 3-year EFS, whereas 2 earlier studies described both survival time
and time to relapse as major endpoints [7,8]. All RCTs included both EFS and OS as efficacy measures
of the trial. Whereas most RCTs focused on localized ES, two had subgroup arms for high-risk and
metastatic disease, which included 277 patients. The mean of the studies’ median follow-up periods
was 6.79 (5.1–8.5) years. Because the median EFS and OS were not reached in 7 and 5 treatment
arms, respectively, analyses regarding median survival were not included in our study. The
radiological response to chemotherapy was not described in any of the studies, and the histological
response was assessed in only 2. Therefore, tumor responses could not be evaluated in the present
study. A significant difference in the HRs of EFS was observed between the control and experimental
arms (HR 0.80, 95% CI 0.68–0.96, \( P = 0.01 \)) (Additional File 2: Figures S1). Meta-analyses of the OS
HRs revealed significantly better survival in the experimental arm than in the standard arm (HR 0.79, 95% CI 0.63–0.98, \( P = 0.03 \)) (Additional File 3: Figures S2). Figure 2 displays forest plots for the treatment effects estimated by hazard ratios (HR) on the 2-year OS and 1-year PFS, TTP, and TTF for each trial.

**Correlations between EFS and OS** The trial-level correlation between HRs for EFS and OS was good (\( R^2 = 0.747, 95\% \text{ CI} 0.531–0.981 \)) (Table 1, Figure 2a). The Spearman’s rank correlation coefficient (\( \rho \)) was 0.683 (95% CI 0.035–0.927, \( P = 0.042 \)). However, the \( R^2 \) for the association between the OS HR and the 1-year EFS was moderate (\( R^2 = 0.348, 95\% \text{ CI} 0.00–0.759; \rho = 0.450, 95\% \text{ CI} -0.305–0.858, P = 0.22 \)) (Table 1, Figure 2b). The correlations between the OS HR and the 3-year EFS (\( R^2 = 0.765, 95\% \text{ CI} 0.545–0.985; \rho = 0.717, 95\% \text{ CI} 0.10–0.936, P = 0.030 \)) and the 5-year EFS (\( R^2 = 0.695, 95\% \text{ CI} 0.423–0.967; \rho = 0.767, 95\% \text{ CI} 0.209–0.948, P = 0.016 \)) were assessed as very good and good, respectively (Table 1, Figures 2c-d).

Similar to what we observed for the 1-year EFS, the correlation between the 1-year OS and the OS HR was poor (\( R^2 = 0.089, 95\% \text{ CI} 0.00–0.408; \rho = 0.214, 95\% \text{ CI} -0.642–0.833, P = 0.64 \)). Meanwhile, the 3-year OS (\( R^2 = 0.831, 95\% \text{ CI} 0.650–1.00; \rho = 0.929, 95\% \text{ CI} 0.584–0.990, P = 0.0025 \)) and the 5-year OS (\( R^2 = 0.809, 95\% \text{ CI} 0.625–0.993; \rho = 0.767, 95\% \text{ CI} 0.209–0.948, P = 0.016 \)) showed very good correlations with OS HR (Table 1, Figures 3a-c).

Further sensitivity analyses were conducted for surrogacy evaluation by removing the treatment arms of the metastatic and high-risk populations. There were 2 RCTs (INT-0091 and EICESS-92) in which metastatic ES was included. In INT-0091, all 120 patients in the metastatic subgroup had metastatic disease [9]. On the other hand, in EICESS-92, the definition of “high-risk” was large localized tumor (≥100 ml) or metastatic disease [10]. Thus, the high-risk subgroup in EICESS-92 included 157 patients with metastatic disease and 335 patients with non-metastatic large localized tumor. After the removal of these subgroups, localized ES analyses revealed an improved correlation between the surrogate endpoints and OS. The correlation between the EFS HR and the OS HR was very good (\( R^2 = 0.818, \)
95% CI 0.625–1.00; \( \rho = 0.929, 95\% \text{ CI } 0.584–0.990, P = 0.0025 \) (Table 2, Figure 4a). The \( R^2 \) for the associations between the OS HR and the 1-year EFS remained moderate \( (R^2 = 0.436, 95\% \text{ CI } 0.00–0.873; \rho = 0.750, 95\% \text{ CI } -0.007–0.961, P = 0.052 \) (Table 2, Figure 4b). The correlations between the OS HR and the 3-year EFS \( (R^2 = 0.807, 95\% \text{ CI } 0.604–1.00; \rho = 0.857, 95\% \text{ CI } 0.294–0.979, P = 0.014 \) and the 5-year EFS \( (R^2 = 0.772, 95\% \text{ CI } 0.537–1.00; \rho = 0.929, 95\% \text{ CI } 0.584–0.990, P = 0.0025 \) were very good (Table 2, Figures 4c-d). The correlation between the 1-year OS and the OS HR remained poor \( (R^2 = 0.136, 95\% \text{ CI } 0.00–0.535; \rho = 0.257, 95\% \text{ CI } -0.701–0.884, P = 0.62) \); however, the 3-year OS \( (R^2 = 0.858, 95\% \text{ CI } 0.693–1.00; \rho = 0.943, 95\% \text{ CI } 0.559–0.994, P = 0.0048 \) and the 5-year OS \( (R^2 = 0.895, 95\% \text{ CI } 0.778–1.00; \rho = 0.929, 95\% \text{ CI } 0.584–0.990, P = 0.0025 \) showed nearly excellent correlations with OS HR (Table 2, Figures 5a-c). When two very old RCTs conducted in the 1970s were further removed from the evaluation of surrogacy, the correlation between HRs for EFS and OS was good, with \( R^2 = 0.519 \) \( (95\% \text{ CI } 0.041–0.997) \) and \( \rho = 0.800 \) \( (95\% \text{ CI } -0.280–0.986) \).

**Discussion**

In RCTs of ES, EFS is often selected as the primary endpoint. EFS has its value, and the advantages of EFS over OS as an endpoint of RCTs are that EFS can be available in a shorter period with a smaller sample size, and subsequently at a lower cost than OS, and that post-progression treatments do not influence EFS. On the other hand, the disadvantages of EFS are that EFS can be affected by the frequency and subjectivity of the surveillance of events, thus including some ambiguity, and that EFS does not always correlate with OS. However, the correlation between EFS and OS has never been verified in ES trials. In this study, data from RCTs of systemic chemotherapy for ES were collected, and the surrogacy of EFS for the true endpoint OS was analyzed using a meta-analytic approach. We identified only 7 relevant trials, reflecting the rare occurrence of the disease. This is the first study to verify the surrogacy of EFS for OS in RCTs of ES. The correlation between the EFS and OS HR in RCTs of newly diagnosed ES was assessed as good, with \( R^2 = 0.747 \) and \( \rho = 0.683 \). Several RCTs used in this analysis included metastatic disease, and not all enrolled patients had localized ES. Although metastatic and localized ES have the same primary treatment strategy, the associated prognoses are
quite different, making it difficult to analyze them together. Therefore, sensitivity analyses were performed with only the treatment arms for cases with localized disease, excluding metastatic and high-risk disease cases, and the correlation between EFS and OS showed an $R^2$ of 0.818 and $\rho$ of 0.929, indicating a very good correlation. The same trend was confirmed by sensitivity analyses of the other time-to-event endpoints, i.e., 1- to 5-year EFS and 1- to 5-year OS. Our results suggest that EFS has the potential to be used as a surrogate for OS in RCTs of newly diagnosed and localized ES. The reason for the better correlation of EFS and OS in localized ES was unknown. This phenomenon needs to be explained by analyzing individual patient data. The average median follow-up time of the included RCTs was 6.79 years. The correlation between 3-year EFS and OS ($R^2 = 0.765, \rho = 0.717$) was comparable to that of the EFS HR and OS, and analyses of the 3-year EFS restricted to localized disease cases further showed a very good correlation with OS ($R^2 = 0.807, \rho = 0.857$). When EFS is selected as the primary endpoint in RCTs of ES, it is considered necessary to conduct follow-up for at least 3 years. In one eligible RCT, 3-year EFS was the primary endpoint [10]. In the European Intergroup Cooperative Ewing's Sarcoma Study-92 trial, the high- and low-risk subgroups included 157 and 4 patients with metastatic disease, respectively. Because the correlation of OS with 3-year EFS tended to be more favorable when examining localized ES, if 3-year EFS is chosen as the primary endpoint, studies limited to localized ES cases may be preferable. In localized lung cancer, a re-analysis of six meta-analyses involving 60 RCTs with 15,071 patients demonstrated that the strength of the correlation between disease-free survival (DFS) and OS in RCTs of adjuvant chemotherapy was $R^2 = 0.92$, which was excellent. For operable lung cancer, DFS is a reliable surrogate endpoint in RCTs performed in adjuvant settings [14]. In RCTs of resectable gastric cancer, the correlation between DFS and OS is also very high ($R^2 = 0.964$), and DFS was concluded to be an acceptable surrogate for OS [15]. On the other hand, the evaluation of the surrogacy of DFS for OS in 22 RCTs of gastro-esophageal cancer demonstrated that DFS did not correlate with OS ($R^2 = 0.27$) [16]. In an analysis of 6 RCTs of adjuvant chemotherapy, which included 12,676 cases of localized colon cancer, the overall correlation between DFS and OS was modest to poor ($R^2 = 0.37$), and DFS was not a good surrogate
for OS in RCTs of stage II colon cancer [17]. Furthermore, in the field of sarcoma, the surrogacy of progression-free survival (PFS) for OS was also investigated using RCTs of advanced soft tissue sarcomas. The correlation between PFS and OS was modest at best; thus the surrogacy of PFS could not be confirmed in the analyses [18, 19]. These observations suggest that confirmation of the correlation between surrogate endpoints and OS is important. Response to chemotherapy may be the surrogate endpoint for which results are obtained most quickly. In RCTs of osteosarcoma, the treatment strategy for changing the postoperative regimen is widely based on the histological response to preoperative chemotherapy [20,21]. Conversely, none of the ES trials selected histological response as an endpoint. Histological response is a significant predictor of favorable outcomes in ES [22]. Nevertheless, only two studies examined histological responses using Huvos grading criteria in the excised tumor tissue [10,13]. Because the strategy of changing the postoperative regimen based on histological response has not been established for ES [2], histological response may not necessarily be investigated in ES trials.

Our study had some limitations. 1) It was based on published, and not individual data. 2) Only 7 RCTs of ES were eligible; thus, the number of RCTs evaluated in the surrogacy analysis was minimal. 3) Definitions and start dates of the time-to-event endpoints varied across trials. 4) The study phase was not clarified in any RCT, and there was no description of post-protocol treatments. An ITT analysis description was observed in only 2 studies, suggesting that the quality of the studies was not excellent. 5) Metastatic disease was included in several trials, and not all studies focused purely on localized ES. In total, 281 (7.8%) of 3,612 patients had metastatic disease. This limits the ability to draw a conclusion from the study results. 6) The study period for each RCT was any period from 1973 to 2010. This might increase the heterogeneity among the RCTs included in this study.

Conclusions
As a surrogate endpoint in RCTs of newly diagnosed ES, EFS showed a good correlation with OS, and the correlation was stronger in cases of a localized disease. EFS may be a useful surrogate endpoint in RCTs of ES, and the surrogacy of EFS is worth verifying using individual patient data in the future.

Abbreviations
Ethics approval and consent to participate

Not applicable.

Consent for publication

Not applicable.

Availability of data and materials

The datasets generated and analyzed in the current study are available from the corresponding author on reasonable request.

Competing interests

The authors declare that they have no competing interests.

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Authors' contributions

KT and HT conceived the study concept and design. KT, MK, TI, and II performed the systematic search of the literature and extraction of data. KT, MK, TI, II, and HT performed statistical analysis, interpretation of data, and drafting and revision of the manuscript. All authors have read and approved the final manuscript.

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Tables
### Table 1. Correlations between surrogate endpoints and OS

| Surrogate endpoint | $R^2$ (95% CI) | $\rho$ (95% CI) |
|--------------------|----------------|-----------------|
| EFS                | 0.747 (0.513–0.981) | 0.683 (0.035–0.92) |
| 1-year EFS         | 0.348 (0.00–0.759) | 0.450 (-0.305–0.8) |
| 3-year EFS         | 0.765 (0.545–0.985) | 0.717 (0.10–0.93) |
| 5-year EFS         | 0.695 (0.423–0.967) | 0.767 (0.209–0.94) |
| 1-year OS          | 0.089 (0.00-0.408) | 0.214 (-0.642-0.8) |
| 3-year OS          | 0.831 (0.650-1.00) | 0.929 (0.584-0.95) |
| 5-year OS          | 0.809 (0.625-0.993) | 0.767 (0.209-0.94) |

Abbreviations: CI, confidence interval; OS, overall survival; EFS, event-free survival.

### Table 2. Correlations between surrogate endpoints and OS in localized Ewing sarcoma

| Surrogate endpoint | $R^2$ (95% CI) | $\rho$ (95% CI) |
|--------------------|----------------|-----------------|
| EFS                | 0.818 (0.625-1.00) | 0.929 (0.584-0.95) |
| 1-year EFS         | 0.436 (0.00-0.873) | 0.750 (-0.007-0.9) |
| 3-year EFS         | 0.807 (0.604-1.00) | 0.857 (0.294-0.97) |
| 5-year EFS         | 0.772 (0.537-1.00) | 0.929 (0.584-0.95) |
| 1-year OS          | 0.136 (0.00-0.535) | 0.257 (-0.701-0.8) |
| 3-year OS          | 0.858 (0.693-1.00) | 0.943 (0.559-0.95) |
| 5-year OS          | 0.895 (0.778-1.00) | 0.929 (0.584-0.95) |
Abbreviations: CI, confidence interval; OS, overall survival; EFS, event-free survival.

Additional File Legends

**Additional File 1: Table S1.** Detailed description of RCTs. RCT, randomized controlled trial; ITT, intention-to-treat; OS, overall survival; EFS, event-free survival; DFS, disease-free survival; VCR, vincristine; ActD, actinomycin D; CPA, cyclophosphamide; ADM (DOX), doxorubicin; IFM, ifosfamide; ETP, etoposide; VAC, VCR+ActD+CPA; VACA, VCR+ActD+CPA+ADM; VAIA, VCR+ActD+IFM+ADM; VAI, VCR+ActD+IFM; VDC, VCR+doxorubicin+CPA; IE, IFM+ETP; EVAIA, ETP+VCR+ActD+IFM+ADM.

**Additional File 2: Figure S1.** Forest plot of EFS with standard vs experimental chemotherapy. CI, confidence interval; EFS, event-free survival; HR, high risk; IV, inverse variance; Meta, metastatic disease; N-meta, non-metastatic disease; SE, standard error; SR, standard risk.

**Additional File 3: Figure S2.** Forest plot of OS with standard vs experimental chemotherapy. CI, confidence interval; HR, high risk; IV, inverse variance; Meta, metastatic disease; N-meta, non-metastatic disease; OS, overall survival; SE, standard error; SR, standard risk.

**Additional File 4: PRISMA flow diagram.

Figures
Figure 1

PRISMA flow diagram. PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses; RCT, randomized controlled trial.
Figure 2

Correlation of EFS with OS HR for Ewing sarcoma. Correlation of (a) EFS HR, (b) 1-year EFS, (c) 3-year EFS, and (d) 5-year EFS. HR, hazard ratio; OR, odds ratio; OS, overall survival; EFS, event-free survival.
Correlation of OS HR with (a) 1-year OS, (b) 3-year OS, and (c) 5-year OS in Ewing sarcoma.

HR, hazard ratio; OR, odds ratio; OS, overall survival.
Correlation of EFS with OS for localized Ewing sarcoma. Correlation of (a) EFS HR, (b) 1-year EFS, (c) 3-year EFS, and (d) 5-year EFS. HR, hazard ratio; OR, odds ratio; OS, overall survival; EFS, event-free survival.
Figure 5
Correlation of (a) 1-year OS, (b) 3-year OS, and (c) 5-year OS with OS HR for localized Ewing sarcoma. HR, hazard ratio; OR, odds ratio; OS, overall survival.

Supplementary Files
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