Dear Editor,

Diffuse large B-cell lymphoma (DLBCL) is the most frequently diagnosed non-Hodgkin lymphoma (NHL) [1]. Rituximab, cyclophosphamide, doxorubicin, vincristine and prednisone (R-CHOP) is the standard treatment for DLBCL. Nevertheless, doxorubicin, like all anthracyclines, is associated with dose-dependent cardiotoxicity [2]. Particularly in patients with congestive heart failure and patients previously exposed to anthracyclines, the use of doxorubicin is contraindicated. In the Netherlands, doxorubicin is most frequently replaced by etoposide (R-CEOP) for anthracyclines-ineligible patients, but efficacy data regarding this regimen are scarce. Recently, two population-based studies from Canada reported on the outcome of R-CEOP. R-CEOP was feasible, although the two studies showed conflicting results [3, 4]. While one study showed inferior outcome of patients treated with R-CEOP, the other observed no difference in disease-specific survival (DSS) between R-CEOP and R-CHOP. Randomized clinical trials (RCT) would be needed to evaluate the efficacy of R-CEOP unbiased, compared to R-CHOP. However, RCTs among anthracyclines-ineligible patients would be unethical, as patients randomized to the doxorubicin-group would experience severe cardiotoxicity due to their cardiac dysfunction. Therefore, propensity-score-matching using population-based data is needed. The aim of this population-based study was to determine the efficacy of R-CEOP in anthracycline-ineligible patients with DLBCL.

We identified all patients ≥18 years diagnosed with DLBCL who received at least one cycle of R-CHOP or R-CEOP between 2014 and 2018, using the Netherlands Cancer Registry (NCR) [5]. Information on patient characteristics and treatment is routinely recorded in the NCR by trained registrars of the NCR through retrospective medical records review. Since 2014, the Lugano classification has been used by the physicians for response evaluation in the Netherlands. Information on the last known vital status for all patients (i.e. alive, death, or emigration) is obtained through annual linkage with the Nationwide Population Registries Network that holds vital statistics on all residents in the Netherlands. Patients alive were censored on February 1st, 2021.

The primary endpoints were progression-free survival (PFS), overall survival (OS) and relative survival (RS). OS was defined as the time between diagnosis and death from any cause, and PFS as the time between diagnosis and tumor progression or death, whichever occurred first. RS was defined as the ratio of the overall survival (OS) of the patient cohort to the expected OS of an equivalent group from the general population, matched to the patients by age, sex, and calendar year. As such, RS reflects the overall excess mortality associated with a DLBCL diagnosis, thereby estimating DSS in the absence of information on the cause of death. The secondary endpoint was overall response rate (ORR), defined as a response of partial remission of the disease or better.

Patients were assigned to the R-CEOP group when they had received 50% or more of their cycles with etoposide instead of doxorubicin. Patients with R-CEOP were propensity-score-matched with patients with R-CHOP in a 1:4 ratio, including age, sex, Ann Arbor stage and International Prognostic Index (IPI) score to account for baseline differences. The log-rank test was used to evaluate differences in survival distributions. Univariable and multivariable proportional hazards regression analyses were performed to assess the impact of treatment on survival after adjustment of sex, age, Ann Arbor stage, IPI score and number of cycles, thereby calculating hazard ratios (HR) and corresponding 95% confidence intervals (95% CI). The Privacy Review Board of the NCR approved the use of anonymous data for this study.

Between 2014 and 2018, 87 DLBCL patients with R-CEOP were matched to 333 DLBCL patients treated with R-CHOP and were included in our study. Median age of the total group of patients was 74 years (range, 39–91 years) and 67% had an advanced stage (Ann Arbor ≥ 3). A total of 42 (48%) patients had high-risk disease (IPI score ≥ 3). Of the 87 patients treated with R-CEOP, 27 patients (31%) had a prior malignancy, as compared to 73 patients (22%) treated with R-CHOP. Baseline characteristics according to R-CEOP and R-CHOP are presented in Table 1.

The median number of cycles administered was 6 (range, 1–8) for both R-CEOP and R-CHOP regimens. Of the 87 patients treated with R-CEOP, 67 (77%) received R-CEOP only, and 20 patients (23%) received both R-CEOP and R-CHOP, of whom the median number of R-CEOP cycles was 5 (range, 3–8) and of R-CHOP 2 (range, 1–4). Among the patients treated with R-CEOP, 13 patients (15%) received subsequent radiotherapy, as compared to 48 patients (14%) in the R-CHOP group. Three patients (3%) treated with R-CEOP received CNS-prophylaxis, as compared to 21 patients (6%) treated with R-CHOP.

The ORR was not significantly different between patients treated with R-CEOP and patients treated with R-CHOP (75% vs. 83%, respectively; p = 0.15; Fig. 1). The CR rates in the R-CEOP and R-CHOP groups were 61% and 72%, respectively (p = 0.21).

The median follow-up was 38 months (interquartile range [IQR], 16–58 months). The 4-year PFS was inferior for patients treated with R-CEOP (44%; 95% CI, 33%–55%) as compared to R-CHOP (58%; 95% CI, 52%–63%; p = 0.03; Fig. 2A). The 4-year OS was 48% (95% CI, 36%–59%) in the R-CEOP group vs. 62% (95% CI, 57%–68%; p = 0.05; Fig. 2B) in the R-CHOP group. However, the 4-year RS was not statistically significantly different for patients treated with R-CEOP (54%; 95% CI, 41%–67%) as compared to R-CHOP (67%; 95% CI, 59%–75%).
Best observed response in patients with diffuse large B-cell lymphoma treated with R-CEOP and R-CHOP. Stacked bar graph depicting best response showing no significant difference in overall response in patients treated with RCEOPs compared to patients treated with R-CHOP ($p = 0.15$).
Fig. 2  Survival analysis. A Progression-free survival (PFS) of patients with diffuse large B-cell lymphoma (DLBCL) treated with R-CEOP and R-CHOP. Kaplan–Meier curves showing significant inferior 4-year PFS in patients with DLBCL treated with R-CEOP ($p = 0.03$). B Overall survival (OS) of patients with diffuse large B-cell lymphoma (DLBCL) treated with R-CEOP and R-CHOP. Kaplan–Meier curves showing a significant inferior 4-year OS in patients with DLBCL treated with R-CEOP ($p = 0.05$). C Relative survival (RS) of patients with diffuse large B-cell lymphoma (DLBCL) treated with R-CEOP and R-CHOP. Kaplan–Meier curves showing no significant difference in 4-year RS in patients with DLBCL treated with R-CEOP ($p = 0.77$).

Table 2. Results of the uni- and multivariable Cox regression analysis on progression-free survival and overall survival for patients with diffuse large B-cell lymphoma treated with R-CEOP versus R-CHOP.

| Covariate              | Univariable PFS | Multivariable PFS | Univariable OS | Multivariable OS |
|------------------------|------------------|-------------------|----------------|------------------|
|                        | HR 95% CI | P-valueb          | HR 95% CI | P-valueb          | HR 95% CI | P-valueb          | HR 95% CI | P-valueb          |
| R-CEOP                 | No      | 1 Reference 1     | Yes       | 1.43 1.02–2.00 0.04 | 1.43 1.00–2.03 0.05 | 1.55 1.11–2.17 0.01 | 1.58 1.11–2.25 0.01 |
|                        | Sex     | Female 1 Reference | Male      | 1.54 1.12–2.10 <0.01 | 1.48 1.07–2.06 0.02 | 1.50 1.09–2.06 0.01 | 1.40 1.00–1.95 0.05 |
|                        | Age     | 18–60 1 Reference | ≥61       | 1.94 1.22–3.09 <0.01 | 2.30 1.37–3.86 <0.01 | 2.36 1.76–3.17 <0.01 | 2.95 2.16–4.01 <0.01 |
|                        | Ann arbor stage | I, II 1 Reference 1 Reference | III, IV 2.17 1.51–3.12 <0.01 | 2.13 1.45–3.12 <0.01 | 2.03 1.25–3.30 <0.01 | 1.87 1.11–3.16 0.02 |
|                        | IPI score | 0–2 1 Reference 1 Reference | 3–5 2.12 1.57–2.86 <0.01 | 2.28 1.65–3.14 <0.01 | 1.79 1.20–2.67 <0.01 | 2.16 1.39–3.36 0.01 |
|                        | Number of cycles | ≥6 cycles 1 Reference 1 Reference | <6 cycles 2.36 1.76–3.17 <0.01 | 2.95 2.16–4.01 <0.01 | 3.30 2.42–4.49 <0.01 | 4.25 3.08–5.87 <0.01 |

Abbreviations: IPI International Prognostic Index, CI confidence interval.

Each covariate is simultaneously adjusted for all other covariates in the table.

b$P$-values are compared with the reference category.
it is most likely that the difference in OS is due to comorbidities. R-CEOP is the treatment of choice for anthracycline-ineligible patients.

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**DATA AVAILABILITY**

The data that support the findings of this study are available via The Netherlands Comprehensive Cancer Organisation. These data are not publicly available, and restrictions apply to the availability of the data used for the current study. However, these data are available upon reasonable request and with permission of The Netherlands Comprehensive Cancer Organisation.

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**AUTHOR CONTRIBUTIONS**

MN designed the study; MB collected the data; MB, MD, and DA-S analyzed the data; MN, MB, MD, and DA-S interpreted the results and drafted the manuscript. All authors read and approved the final version of the manuscript.

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**COMPETING INTERESTS**

The authors declare no competing interests.

**ETHICS APPROVAL**

According to the Central Committee on Research involving Human Subjects (CCMO), this type of observational study does not require approval from an ethics committee in the Netherlands. The Privacy Review Board of the NCR approved the use of anonymous data for this study.

**ADDITIONAL INFORMATION**

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