Anthraclycline cardiotoxicity is traditionally described as type 1 cardiotoxicity that features cardiomyocyte death resulting in irreversible injury. This is in contrast to type 2 cardiotoxicity, associated with trastuzumab, that is characterized by cardiomyocyte dysfunction, rather than cell death, and is therefore felt to be reversible. In the study by Cardinale et al, initiation of cardioprotective medications promptly after prompt initiation of enalapril, either alone or in conjunction with a β-blocker, with severity of postchemotherapy LV dysfunction and New York Heart Association functional class being the best predictors of recovery. The study findings suggest that anthraclycline cardiotoxicity represents a continuum that begins with subclinical myocardial cell injury, followed by an early asymptomatic decline in LVEF that can progress to symptomatic heart failure, if left untreated (Figure). The few cases that presented late occurred in patients with confounding risks for the development of cardiomyopathy, raising questions about the involvement of anthraclyclines or suggesting a double-hit phenomenon in patients with a myocardium made vulnerable by previous anthraclycline treatment. These findings challenge the existing dogma that anthraclycline cardiotoxicity represents irreversible myocardial injury that can present as either acute, early-onset chronic progressive, or late-onset chronic progressive cardiotoxicity. The prospective and nonselective, symptom-independent approach to LVEF surveillance in the current study would suggest that late-onset anthraclycline cardiotoxicity likely reflects the timing of detection, rather than the timing of the occurrence of cardiotoxicity. These findings have significant implications for the surveillance and management of anthraclycline cardiotoxicity.

Although the need for cardiac monitoring of asymptomatic anthraclycline-treated adult patients is generally acknowledged, existing guidelines offer no clear consensus regarding the timing or duration of such surveillance. One proposed schedule includes measurement of LVEF at 6 months following treatment, annually for 2 to 3 years thereafter, and then at 3- to 5-year intervals for life, with more frequent monitoring in high-risk patients. The cost implications of periodic lifetime screening following anthraclycline exposure are unknown, but daunting when one considers the growing population at risk. Indeed, rationing of similar lifelong surveillance schedules used in childhood cancer survivors exposed to anthraclyclines has been suggested in an effort to improve the efficacy and cost-efficiency of screening. Cardinale and colleagues now present evidence to suggest that, in adults with cancer, cardiotoxicity occurs almost exclusively within the first year after completing anthraclycline treatment. Focusing cardiac screening to this high-risk period could significantly improve cost-effectiveness, and physician and patient acceptance and adherence, as well.

The opinions expressed in this article are not necessarily those of the editors or of the American Heart Association.

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the detection of cardiotoxicity was associated with recovery in 82% of patients over a mean period of 8±5 months. These data challenge the concept of type I cardiotoxicity and indicate significant potential for reversibility with early detection and treatment, providing further support for periodic surveillance early after anthracycline therapy. It is interesting and intuitive to note that the mean LVEF before the initiation of heart failure treatment was significantly lower among patients without LVEF recovery. Furthermore, patients without LVEF recovery had a higher incidence of adverse cardiac events. These observations also support the concept of a spectrum of cardiotoxicity in which the likelihood of reversibility reduces as LVEF worsens.

Recent studies have investigated the utility of prophylactic angiotensin-converting enzyme inhibitor and β-blocker use in anthracycline-treated patients. Combined prophylactic treatment with enalapril and carvedilol achieved a lower incidence of death, heart failure, or a final LVEF of <45% in comparison with controls (24.4% versus 6.7%, \( P = 0.02 \)) in a randomized, placebo-controlled trial of 90 patients embarking on predominantly anthracycline-based chemotherapy.\(^\text{10}\) In earlier work by Cardinale and colleagues, troponin elevation within 72 hours of high-dose chemotherapy was used to randomly assign 114 patients to an enalapril-treated or control cohort; the incidence of an absolute decrease in LVEF of >10% from baseline and below the lower limit of normal was significantly higher in control subjects (43% versus 0%, \( P < 0.001 \)). In patients with established LV dysfunction, time to heart failure treatment is an independent predictor of incomplete LV recovery,\(^\text{15}\) suggesting that delays in the initiation of cardioprotective medications should be minimized. Universal prophylaxis with cardioprotective agents, although seemingly effective, commits patients without cardiotoxicity to treatment, which is particularly undesirable in the context of frequent polypharmacy and variable blood pressures during active cancer therapy. Thus, screening and selection of patients for prompt initiation of heart failure therapy based on objective evidence of myocardial injury (eg, troponin elevation, decrease in global longitudinal strain, or decrease in LVEF), with or without symptoms, represents a more clinically reasonable strategy. The high rate of reversible cardiotoxicity in the current study by Cardinale et al\(^\text{2}\) should be validated using screening algorithms that advocate the use of biomarkers and echocardiographic parameters of early myocardial injury to guide therapy.\(^\text{16}\)

In tandem with the introduction and uptitration of cardioprotective agents, the clinical response to cardiotoxicity during chemotherapy often includes a reassessment of chemotherapy regimens, which may result in interruption, dose reduction, or substitution of cancer treatments. Cardinale and colleagues do not describe the breakdown of oncological responses, if any, that accompanied the detection of cardiotoxicity and do not explore associations with oncology end points. These data would be particularly interesting given that 97% of deaths (\( n = 792 \)) were attributable to tumor-related causes.\(^\text{2}\) A fundamental principle governing cardio-oncology practice is that reducing the risk of cardiovascular toxicity must be balanced against maintaining the efficacy of cancer treatment. To date, there are no data on the effects of prospective cardiac monitoring during anthracycline treatment on oncology outcomes. It is imperative that cardiac-monitoring recommendations and downstream reactive changes in cancer treatments do not cause more harm than good.

Despite the incremental contribution of the study by Cardinale et al, there are several unresolved questions surrounding anthracycline cardiotoxicity. In this study, all patients who manifested sufficient LV dysfunction were promptly treated with heart failure medications, irrespective of symptoms. However, the natural history of anthracycline-induced LV dysfunction in terms of spontaneous recovery and cardiovascular outcomes remains uncertain. Similarly, the clinical relevance of troponin elevations or reductions in global longitudinal strain that can precede overt deteriorations in LVEF and the optimal management of these abnormalities require further investigation. Not all subclinical cardiotoxicity progresses to overt LV
dysfunction or warrants intervention, and the challenge lies in distinguishing irrelevant cardiac effects from prognostically meaningful anthracycline toxicity. In the current study, 82% of patients treated for cardiotoxicity demonstrated some recovery, with LVEF returning to baseline values in 11%. Cardiac medications were continued throughout the duration of this study, but the appropriate duration of treatment remains uncertain. Whether cardioprotective agents can be safely weaned or discontinued in patients who achieve full recovery is unclear. Younger and asymptomatic patients may be particularly reluctant to commit to long-term cardioprotective medications in the absence of supportive data. Last, it is uncertain whether the findings of this study are applicable to childhood cancer survivors of anthracycline therapy. A previous randomized trial of enalapril in 135 pediatric survivors with evidence of asymptomatic and symptomatic anthracycline cardiotoxicity, treated at least 2 years after completion of cancer therapy, did not show a benefit in cardiac function. This may suggest divergent pathophysiologies in children and adults exposed to anthracyclines or may reflect a lack of efficacy attributable to the delayed institution of heart failure therapy.

Clinical decision making in cardio-oncology is often limited by a dearth of long-term prospective studies and robust, evidence-based guidelines. The American Society of Clinical Oncology–Cardio-Oncology Survivorship Expert Panel deemed evidence identified from a systematic review of the literature insufficient to support a practice guideline to direct screening for chemotherapy- and radiation therapy–induced cardiac effects. This panel highlights the relative lack of prospective studies in comparison with cross-sectional or retrospective studies in this field. As such, existing guidelines are based on consensus rather than evidence and differences between guidelines have hindered effective implementation of recommendations. This has contributed to significant practice variation in surveillance for and management of anthracycline cardiotoxicity, and in other areas of cardio-oncology, as well. This study by Cardinale and colleagues offers evidence that could inform guidelines and lead to greater standardized cardiac care of patients exposed to anthracyclines. This study will hopefully inspire others to systematically evaluate the role of periodic noninvasive testing for cardiac dysfunction and the role of treatment to prevent cardiac disease in asymptomatic cancer patients receiving other potentially cardiotoxic treatments such as tyrosine kinase inhibitors, proteasome inhibitors, and other novel agents. Cardio-oncologists have a responsibility to push forward with similar clinical research in an effort to improve the cardiovascular care of a growing population of cancer survivors predisposed to adverse cardiovascular outcomes.

Disclosures

None.

References


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