Anti-HER2 therapy-associated cardiotoxicity in breast cancer patients: analysis of real-world data from a UK cancer centre

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Introduction
Cardiac adverse events are a recognised toxicity for patients with breast cancer receiving anti-human epidermal growth factor receptor-2 (HER2) therapy.

Large phase III clinical trials have shown an incidence of left ventricular ejection fraction (LVEF) decrease >10% of 7.1–18.6% and overt heart failure of 1.7–4.1%. Retrospective analyses from large registries such as the Surveillance, Epidemiology and End Results (SEER) and Cancer Research Network have reported an even higher incidence of heart failure at 41.9% and 20.1%, respectively.

We conducted a retrospective analysis to evaluate the incidence of cardiac adverse events for patients receiving anti-HER2 therapy at St Luke's Cancer Centre (SLCC) and Ashford and St Peter's Hospital (ASPH). The aims of the study were to provide real-world data on cardiotoxicity with anti-HER2 therapy and to evaluate adherence to the cardiac surveillance protocol.

Materials and methods
Patients with breast cancer receiving anti-HER2 therapy between June 2018 and June 2019 at SLCC and ASPH were included in the analysis. Data on patient demographics, chemotherapy and anti-HER2 therapy details, and LVEF values were obtained from the systemic anti-cancer therapy electronic prescribing system and patient records.

Results and discussion
One hundred and twenty-three female patients were analysed; the median age was 56 years (range 29–82 years); 25 (23.8%) patients had at least one cardiac comorbidity, with hypertension being the most common; 66% of patients were treated with curative intent; 40% of patients received combined anthracycline–taxane chemotherapy, while 60% received only taxane-based chemotherapy; 55.3% received dual anti-HER therapy with pertuzumab and trastuzumab, while 44.7% were treated with trastuzumab alone. The median number of anti-HER2 therapy doses was 16 (range 3–93). An asymptomatic decrease in LVEF was recorded in 33 patients (26.8%). Of these, 19 patients had a decrease in LVEF >10 points below baseline, six had an LVEF decrease to <50% and eight had both a decrease in LVEF >10 points and <50%. The median time from start of treatment to LVEF decrease was 5 months (range 2–36 months). At the time of reporting, the LVEF of 22 patients (66.7%) had recovered to baseline. There were no cases of symptomatic heart failure. Three patients had delays in their treatment due to the LVEF drop and two stopped treatment early. LVEF decrease was more common in patients receiving palliative treatment (35.7%) vs 22.5%) and anthracycline chemotherapy (28.3% vs 24.6%). No association was found between LVEF decrease and cardiac comorbidity or type of anti-HER2 therapy. Adherence to the cardiac surveillance protocol was 76.7%. The most common deviation from protocol was a delay in the cardiac function assessment at 4 months.

Conclusions
In this real-world analysis of cardiotoxicity with anti-HER2 therapy, the incidence of significant cardiac adverse events and interruption or discontinuation of therapy was very low, confirming the safety of this treatment modality. Asymptomatic decrease in LVEF was associated with palliative treatment intent and anthracycline-based chemotherapy. Adherence to the cardiac surveillance protocol is important, as the majority of LVEF decreases occur within the first 5 months of therapy and are potentially reversible with appropriate interventions.

Conflicts of interest
None declared.

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