Myopericarditis From Bevacizumab/5-Fluorouracil (5-FU) in a Patient Presenting With ST-Elevation Myocardial Infarction (STEMI): A Case Report

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Background

5-fluorouracil (5-FU) and its oral prodrug capecitabine, are extensively utilized with bevacizumab as first-line chemotherapy for metastatic colorectal cancer. These agents are associated with a range of cardiotoxic effects that can be clinically significant.

Case presentation:

We report a case of a Caucasian male in his 70's (Figure 1) who experienced chest discomfort and electrocardiography features of ST-elevation myocardial infarction (Figure 2) on the background of receiving adjuvant 5-FU and bevacizumab, following high anterior resection for node-positive metastatic sigmoid adenocarcinoma. Urgent coronary angiography revealed unobstructed coronary arteries (Figure 3), and cardiac magnetic resonance imaging (Figure 4) confirmed the diagnosis of myopericarditis.

Ethical compliance: Informed consent was obtained from patient.

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Imaging



Figure 1: Medical history timeline

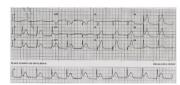


Figure 2: STEMI pre-notification



Figure 3: Mid RCA stenosis with TIMI III flow

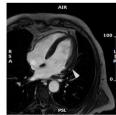


Figure 4: LGE in subepicardial lateral segments

Discussion and Learning Points

- 1. The degradation product of 5-FU (α -fluoro- β -alanine) is converted into fluorocitrate, which disrupts the citric acid cycle and results in the accumulation of citrate, the subsequent depletion of high-energy ATP and ischaemia¹
- The Heart Failure Association—International Cardio-Oncology Society (HFA-ICOS) tool can identify high-risk patients requiring close followup²
- Pharmacokinetic-guided dosing, rather than body surface area adjustment, has been shown to reduce 5-FU toxicity and improve survival²
- 4. For those who develop cardiotoxicity, rechallenge may be cautiously attempted with vasodilator prophylaxis (e.g., diltiazem and isosorbide mononitrate) and close telemetry²
- 5. CMR can distinguish myopericarditis from infarction or Takotsubo cardiomyopathy

Conclusion:

The temporal association with chemotherapy, in the absence of other identifiable causes, strongly implicated 5-FU as the causative agent, with a possible contributory role of coronary vasospasm.

Reference:

- Matsubara I, et al.; Cardiotoxic effects of 5-fluorouracil in the guinea pig. JpnJPharmacol
- Gamelin E. Et al.; Individual Fluorouracil Dose Adjustment Based on Pharmacokinetic Follow-Up Compared With Conventional Dosage Results of a Multicenter Randomized Trial of Patients With Metastatic Colorectal Cancer. JCO 26:2099–2105