Capecitabine-Induced Coronary Vasospasm

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Case Report

Capecitabine-Induced Coronary Vasospasm

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Keywords
Capecitabine · 5-Fluorouracil · Chest pain · Coronary artery disease · Breast cancer

Abstract
Capecitabine, an oral prodrug of 5-fluorouracil (5-FU), is approved for early-stage and advanced colorectal cancer and metastatic breast cancer. Cardiotoxicity of 5-FU is well described in the literature. However, cardiac adverse effects of capecitabine are poorly described. We report a case of coronary vasospasm induced by capecitabine. A 41-year-old female with metastatic breast cancer presented with chest pain 3 days after starting capecitabine. The chest pain was relieved by rest and exacerbated by exertion. Her physical examination was unremarkable except for a rapid heart rate of 100 bpm. Electrocardiogram test showed no acute ischemic changes. Troponin tests were negative. CT angiography of the chest was negative for acute pulmonary embolism. An echocardiogram showed a left ventricular ejection fraction of 60% without any wall motion abnormalities. The chest pain resolved with aspirin and analgesic use. She was discharged following an inconclusive cardiac workup. Further use of capecitabine was discontinued.

Introduction
Capecitabine (Xeloda) is an oral fluoropyrimidine antimetabolite, which is converted by thymidine phosphorylase to 5-fluorouracil (5-FU). This enzyme is preferentially expressed
by tumor cells rendering capecitabine highly selective for sensitive tumor cells [1]. It is en-
listed in the WHO List of Essential Medicines and is currently approved for early-stage and
advanced colorectal cancer and metastatic breast cancer. Cardiotoxicity with fluoropyrimi-
dine-based treatment is occasionally observed, and could even be fatal [2, 3]. Although 5-FU
is known to cause cardiotoxicity, capecitabine-induced cardiac side effects are poorly de-
scribed in the literature. Herein, we report a case of coronary vasospasm induced by capeci-
tabine in a patient with no pre-existing coronary artery disease (CAD) or risk factors.

Case Presentation

A 41-year-old female with metastatic breast cancer presented to the emergency de-
partment with chest pain of 1 day duration. The chest pain started gradually, and was re-
lieved by rest and exacerbated by exertion. She was started on capecitabine therapy 3 days
before the onset of chest pain. She denied shortness of breath, nausea, vomiting, diaphoresis,
fever, syncope, heartburn, and peripheral edema. The patient did not have a history of CAD
or had known risk factors for CAD. She denied the use of cocaine. The patient did not de-
scribe symptoms suggestive of panic attack. On examination, her heart rate was 100 bpm
and regular. Her physical examination was otherwise unremarkable.

Five years before the current event, she underwent mastectomy and axillary lymph node
dissection for a stage III (pT2N3) breast cancer. The tumor was ER+, PR+, HER2+, and
she received adjuvant chemotherapy and trastuzumab, followed by tamoxifen. She was di-
agnosed with metastatic disease in the liver in 2014, and was started on trastuzumab, pert-
tuzumab, and docetaxel. Fifteen months later, she had disease progression in the liver and
was switched to trastuzumab-emtansine. Nine months later, she progressed again, and ther-
apy was changed to lapatinib and capecitabine (1,500 mg b.i.d.). Three days after starting
capecitabine, she experienced acute-onset chest pain.

Electrocardiogram (ECG) testing showed normal sinus rhythm and no changes sugge-
sitive of coronary ischemia. Cardiac enzymes were negative. Complete blood count and chem-
istry panel were within normal range. CT angiography of the chest did not reveal any evi-
dence of pulmonary embolism. A transthoracic echocardiogram documented normal cardiac
function and a left ventricular ejection fraction of 60%. Her chest pain resolved after aspirin
and analgesics were administered upon admission in the ER. A cardiology opinion
was sought for possible angiography, but was not undertaken because of the patient’s improving
condition. She was discharged following complete resolution of chest pain. Oncology was
consulted during the hospitalization, and capecitabine therapy was discontinued. Since then,
she has had no complaints of chest pain. Repeat echocardiogram was normal as well.

Discussion

The selective anti-tumor action of capecitabine increases its tolerability. However, capeci-
tabine and its parental compound may induce cardiotoxicity, depending on the dose, card-
iac comorbidity and schedule of chemotherapy [4]. Capecitabine cardiotoxicity is thought to
occur through the action of 5-FU on the endothelium resulting in production of endothelin-1
and subsequent coronary vasospasm [5].

Patients present with symptoms similar to variant angina, with chest pain at rest and
may or may not have ECG changes of ischemia. Exercise-induced ECG changes have been
observed with capecitabine use [6]. In fact, Goldsmith et al. [7] reported exercise-induced global myocardial ischemia in a patient with normal coronary arteries and left ventricular function who was on capecitabine for recurrent breast cancer.

With regard to onset of chest pain with capecitabine therapy, Wijesinghe et al. [8] reported an acute coronary syndrome in a patient who had been on capecitabine for only 2 days. Goldsmith et al. [7] reported a patient who experienced symptoms due to coronary vasospasm a few days after starting capecitabine. Our patient presented with chest pain 3 days after taking capecitabine. Depending on the dose, cardiac side effects can occur within 24 h after taking the drug [9].

As for our patient, the resolution of chest pain, normal ECG and troponins ruled out myocardial infarction and dysrhythmias. The recording of ST segment changes on ECG depends on the presence of symptoms at the time of assessment. Our patient’s chest pain subsided before ECG was recorded that could have possibly led to the absence of changes suggestive of coronary vasospasm. The normal echocardiogram excluded direct damage to the myocardium or coronary vessels by capecitabine metabolite. It was thought that a coronary angiography and a stress test are not necessary for her management, and therefore it was not obtained.

The cardiotoxicity of fluoropyrimidines manifests as vasospasm, hypertension, ventricular arrhythmias, cardiogenic shock, and even cardiac arrest [2, 5, 10]. Also, there is a higher prevalence of capecitabine cardiotoxicity in patient with CAD. This is because thymidine phosphorylase activity is reported to be expressed in atherosclerotic plaques [11]. Our patient did not have CAD. Of note, 5-FU-mediated cardiotoxicity can occur in patients with no pre-existing CAD [5–7, 12].

In summary, our patient had capecitabine-induced coronary vasospasm in the absence of pre-existing CAD. Coronary vasospasm is reversible with discontinuation of capecitabine. Early recognition of this complication of treatment will help prevent adverse events. Further use of capecitabine must be discontinued to avoid risk for cardiotoxicity.

Statement of Ethics
The authors have no ethical conflicts to disclose.

Disclosure Statement
The authors have no conflicts of interest to disclose.

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