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Development of stress-induced cardiomyopathy after cytoreductive surgery and hyperthermic intraperitoneal chemotherapy

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Abstract

Pseudomyxoma Peritonei, a massive mucinous peritoneal collection due to a rare epithelial neoplasm, can be effectively treated with Cytoreductive Surgery and Hyperthermic Intraperitoneal Chemotherapy (CRS-HIPEC).

A 43-year-old female, previously treated for mucinous ovarian carcinoma with CRS-HIPEC, and total abdominal hysterectomy and bilateral salpingo-oophorectomy, presented with new-onset abdominal distension and early satiety. She was diagnosed with Pseudomyxoma Peritonei. After 48 hours of treatment with CRS-HIPEC, she presented haemodynamically unstable with acute chest pain. Electrocardiogram showed broad complex tachycardia with ST depression in leads V3-6. Severe systolic dysfunction with Ejection Fraction (EF) of 20% along with severe pulmonary hypertension, visualized on Echocardiography. A diagnosis of Stress-induced Cardiomyopathy was established using InterTAK Diagnostic Score.

Patients with CRS-HIPEC have presented with Stress-induced Cardiomyopathy. However, no specific relation between the two has been established. This case report discusses Stress-induced Cardiomyopathy as a complication of CRS-HIPEC.

Keywords: Stress-induced; cardiomyopathy; Cytoreduction; Chemotherapy
Introduction

Stress-induced Cardiomyopathy, also known as Takotsubo Cardiomyopathy Syndrome (TTS) is defined as temporary left ventricular dysfunction due to stress and/or surgery.\(^{(1)}\) Pseudomyxoma Peritonei (PMP), a rare clinical condition which occurs in 1–2 people in a million\(^{(2)}\) is characterized by massive amounts of mucous in the peritoneum mostly due to a tumour of the appendix.\(^{(3)}\) Despite the slow spread, the long term survival rate is 50% and 10%-30% at 5 and 10 years, respectively.\(^{(4)}\) The gold standard treatment of PMP is Cytoreductive Surgery and Hyperthermic Intraperitoneal Chemotherapy (CRS-HIPEC)\(^{(3)}\) as it not only improves the quality of life but has been shown to have better survival rates.\(^{(3)}\) A retrospective multicentre series of 2,298 patients, showed a 10-year survival rate of 63% after CRS-HIPEC.\(^{(5)}\)

However, this treatment can induce physiological and potentially pathological changes causing stress-induced Cardiomyopathy which may lead to postoperative morbidity and mortality. We present the case of a 43-yaer-old female, who developed stress-induced Cardiomyopathy after undergoing CRS-HIPEC for PMP. this case report highlights that stress-induced Cardiomyopathy is a complication of CRS-HIPEC.

Case Report

A 43 years old female presented at Shifa International Hospital, Islamabad on 3th March 2019 with complaints of abdominal swelling and early satiety. Upon taking her medical and surgical history, she had experienced similar symptoms in the past and was diagnosed with Mucinous Ovarian Adenocarcinoma. She underwent elective CRS-HIPEC. She later developed metastasis which were treated by total abdominal hysterectomy and bilateral salpingo-oophorectomy in August 2018. The patient had received four cycles of neoadjuvantive and adjuntive chemotherapy, each with Carboplatin and Paclitaxel. However, she recently started developing notable abdominal swelling and early satiety again.

On evaluation, she was alert and well-oriented individual, with a height of 162 cm and weight 51 kg. Her abdominal CT scan showed findings that were typical of PMP with
the suggestion of rupture of appendiceal mucocoele as the primary cause. However, no pathology was found in her chest. Due to her current disease, she was again offered CRS-HIPEC.

Preoperatively, she had good functional class and her preoperative blood test was within the normal limit except for mild anaemia (Hb = 10.2 g/dl). Her electrocardiogram and echocardiography showed sinus tachycardia and an ejection fraction of 55%.

She underwent CRS which included splenectomy, appendectomy, cholecystectomy, excision of pelvic mass adherent to the bladder, excision and repair of the bladder and peritonectomy. This was followed by HIPEC with Mitomycin 20 mg, at 42 °C for 90 minutes. The intraoperative blood loss was 2000 ml. She was transfused 750 ml of packed red blood cells and six litres of crystalloids and 200 ml of Albumin. She remained stable intraoperatively except for occasional episodes of tachycardia and a need for low-dose norepinephrine infusion to support haemodynamic parameters.

After the surgery, the patient was extubated and shifted to Intensive Care Unit. After forty-eight hours of surgery, she complained of chest pain and developed tachycardia, hypotension, tachypnoea and decrease in oxygen saturation (SpO2: 86% on room air). Her 12-lead ECG showed broad-complex tachycardia and depression of ST segment in leads V3-6. Her echocardiography (Table 1) showed severe systolic dysfunction with EF of 20%, along with severe pulmonary hypertension. Her cardiac enzymes and BNP levels (Table 2) were also raised.

She was started on norepinephrine at 0.04 µg/kg/min. Aspirin and clopidogrel were administered. Enoxaparin 60 mg subcutaneous was given on the following day. Her respiration was supported by Bilevel Positive Airway Pressure. Cardiology consult recommended coronary angiography but was refused by the patient. Her condition improved overtime and she was discharged on the seventh post-operative day. On 2-weeks follow-up visit, her echocardiogram revealed an improvement in ejection fraction to 45%. No other abnormalities were noted, and she was scheduled for regular follow-ups.
Discussion

CRS-HIPEC involves removal of the PMP affected areas, peritoneum and other intra-abdominal organs, followed by perfusion of heated chemotherapy agent at 42-43° C for 1-2 hours directly within the abdominal cavity. Broadly, the procedure includes three stages: Exploration, Cytoreduction/debulking and Chemoperfusion. CRS has minimal residual tumour and cytotoxicity is achieved by delivering chemotherapeutic agents directly into the peritoneal cavity. ⁶)

The morbidity after CRS-HIPEC can be either related to surgery or chemotherapy. Chemotherapy effects wound healing and increases the risks of infectious complications. The surgery-related morbidity includes abscess, fistula, anastomotic leak, postoperative ileus, wound infection, bleeding, thrombosis, pulmonary embolism, pleural effusion, pneumothorax, and cardiac arrhythmias. (⁷) However, there is no case report showing a relationship between CRS-HIPEC and stress cardiomyopathy.

TTS on electrocardiogram may mimic acute coronary syndrome. It may also present with raised myocardial enzymes even with no obstructive coronary artery disease. However, coronary angiography is often normal or typically shows less than 50% luminal stenosis. Most patients recover completely, and the severely reduced left ventricular systolic function recovers over four to eight weeks. (⁸)

Stress-induced Cardiomyopathy is predominant in females. Patients mostly present with chest pain and dyspnoea. However, other more serious signs such as cardiogenic shock, hypotension, arrhythmias, and cardiopulmonary arrest are relatively common. (⁸) TTS is most significantly associated with an identifiable stressor, ranging from an emotional stressful event without a physical component to a physical stressor. Typically, physical stressors involve major surgery, orthopaedic trauma, exacerbation of obstructive airways disease, malignancy, chemotherapy and infections. (⁹)

In TTS, as compared to ACS, initially the troponin values are usually similarly raised, however, the peak values are noticeably lower. There is only slight elevation in creatinine kinase and a remarkable increase in the plasma BNP which continues to rise till approximately 24–48 hours after symptom onset. (¹⁰)
In order to correctly diagnose Stress-induced Cardiomyopathy, the International Takotsubo Registry provides clinicians with the InterTAK Diagnostic Score. According to the scoring system, the predicted probability of TTS increases with the increasing score points of a patient. There is a chance of less than 1%, if the patient has 30 score points. Similarly, patients with 50 points are 18% and patients with more than 70 points are 90% more likely to have TTS.¹⁰

In the present case, the patient had emotional stress, and physical stress, adding up to a score of 62 points indicating more than fifty percent probability of TTS. Although the mechanism of TTS for this patient was uncertain, its occurrence seems to be associated with anaesthesia, surgery, post-operative pain, chemotherapy and excessive stress inflicted by the outcome of this postoperative treatment. This patient had chest pain, dyspnoea, T-wave inversion in leads V3-V5 on ECG, mildly elevated cardiac enzymes and BNP levels, and akinetic mid antero-septum on echocardiography. All of which leads to a diagnosis of stress-induced Cardiomyopathy.

As she had high predictive InterTAK diagnostic score and no pre-operative tests indicating coronary artery disease, coronary angiography to rule out coronary artery disease was not performed. In addition, all the post-operative blood investigations and imaging studies established the diagnosis of stress-induced Cardiomyopathy.

**Conclusion**

This case of a 43-year-old female highlights stress-induced Cardiomyopathy as a complication of the gold standard treatment modality available for PMP. In this case report, the diagnosis of stress-induced Cardiomyopathy was made using the InterTAK Diagnostic Score, and factors related to the presentation, aetiology, associations and management are identified to prevent morbidity and mortality associated with the treatment modality.
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Conflict of interest: None to declare

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References


Abbreviations

ACS: acute coronary syndrome; AR: aortic regurgitation; BNP: Brain natriuretic peptide; CAD: Coronary artery disease; CK-MB: Creatine kinase-MB; CMR: Cardiovascular magnetic resonance imaging; CRP: C-reactive protein; CRS: cytoreductive surgery; CT: computerized tomography; DVT: deep vein thrombosis; ECG: electrocardiogram; Echo: echocardiogram; ESR: erythrocyte sedimentation rate; Hb: Haemoglobin; HIPEC: hyperthermic intraperitoneal chemotherapy; LVOTO: Left ventricular outflow tract obstruction; METs: metabolic equivalent; MR: mitral regurgitation; PMP: Pseudomyxoma peritonei; POD: postoperative day; PR: pulmonary regurgitation; RV: Right Ventricle; RWMA: regional wall motion abnormalities; SPO2: peripheral capillary oxygen saturation; TR: tricuspid regurgitation; TTE: transthoracic echocardiogram; TTS: Takotsubo cardiomyopathy syndrome
**Table 1: Echocardiography reports**

<table>
<thead>
<tr>
<th>Echocardiography reports</th>
<th>Preoperative Echo</th>
<th>EF: 55%, Trace AR, TR, mild mitral valve prolapse with mild MR</th>
</tr>
</thead>
<tbody>
<tr>
<td>ECHO on POD 2</td>
<td>EF = 20%, mid antero-septum is akinetic, severe pulmonary hypertension, mild MR, moderate TR, mild PR, severe cardiomyopathy</td>
<td></td>
</tr>
<tr>
<td>ECHO on POD 4</td>
<td>Normal left ventricular size with severe systolic dysfunction. Estimated EF = 25%. Mid anteroseptum is akinetic.</td>
<td></td>
</tr>
<tr>
<td>Echo on POD 14</td>
<td>EF =40%</td>
<td></td>
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**Table 2: Cardiac enzymes and BNP levels**

<table>
<thead>
<tr>
<th>Cardiac enzymes and BNP levels</th>
<th>Immediately after the event</th>
<th>Six hours after the event</th>
<th>Nine days after the event</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cardiac troponin I</td>
<td>555.9 pg/mL</td>
<td>7397.3 pg/mL</td>
<td>-</td>
</tr>
<tr>
<td>(reference range in females upto 15.6 pg/mL)</td>
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<tr>
<td>Creatine kinase MB (CK-MB)</td>
<td>4.2 ng/mL</td>
<td>30.5 ng/mL</td>
<td>-</td>
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<tr>
<td>(reference range in females up to 3.4 ng/mL)</td>
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<td></td>
<td></td>
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<tr>
<td>Brain natriuretic peptide (BNP)</td>
<td>252.1 pg/mL</td>
<td>-</td>
<td>190 pg/mL</td>
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<td>(reference range &lt;100 pg/mL)</td>
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