An Ovarian Cancer Patient Presenting Heparin-Induced Thrombocytopenia after use of Low Molecular Weight Heparin: A Case Report of Rare Condition

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ABSTRACT

Heparin-induced thrombocytopenia is a rare and life-threatening condition of exposure to heparin. A case of heparin-induced thrombocytopenia due to the low molecular weight heparin was presented. Pulmonary emboli and progressively decreased number of thrombocytes developed during the low molecular weight heparin treatment. For that reason, the heparin-induced thrombocytopenia was diagnosed. The heparin was ceased and fondaparinux treatment initiated. Platelet levels returned to normal limits within six days. The delaying in diagnosis of heparin-induced thrombocytopenia causes serious outcomes. The physician must be careful and keep in mind be developed of this clinical condition in patient under heparin treatment.

Keywords: Heparin-induced thrombocytopenia, Low molecular weight heparin, Ovarian cancer

Introduction

Heparin-induced thrombocytopenia (HIT) is an immune-mediated, life-threatening condition of exposure to heparin (unfractionated heparin; UFH or low molecular weight heparin; LMWH) that occurs in a small percentage of patients exposed, regardless of the dose or administration (1). HIT results from an autoantibody directed against endogenous platelet factor 4 (PF4) in complex with heparin. HIT antibodies remain in blood circulation for several weeks to several months (2). This immune reaction activates platelets and can increase the risk of catastrophic arterial and venous thrombo- sis. HIT is estimated to occur in <0.1 to 5.0% of patients receiving heparin depending on the clinical scenario (1-3). Unfractionated heparin and LMWH are commonly used agents for prophylaxis against venous thromboembolism after surgery and in patients admitted to the hospital (3). A potential advantage of LMWH from clinical and cost perspectives is a lower risk for HIT (4).

The most important risk factors for HIT are; duration of heparin therapy, type of heparin, indication for UFH/LMWH, and sex of the patient (5). Clinically, HIT may lead to asymptomatic or symptomatic thrombocytopenia or manifest as skin lesions at the heparin injection site or with acute systemic reactions like chills, fever, dyspnea, chest pain, bleeding (intracranial/retroperitoneal) or thrombosis like pulmonary embolism, myocardial infarction, cerebrovascular disease, necrosis of extremity.

The following steps should be followed when diagnosing HIT; evidence of longer than 5 days’ heparin exposure, omission of other causes of thrombocytopenia, demonstration of HIT antibodies by functional or antigenic methods (ELISA, serotonin release assay, heparin-induced platelet aggregation test, flow cytology) and elevated platelet counts 4 to 14 days after heparin is discontinued should be observed.

This case report is written to indicate that HIT should always be remembered when a patient’s platelet counts decrease.
after 5-10 days of heparin treatment. Here is described the case of a patient with HIT syndrome due to LMWH use after an ovarian cancer surgery.

**Case Report**

A 74-year-old patient underwent surgery for advanced stage ovarian cancer. Intraoperative frozen section assessment of omental cake was reported as malignant epithelial tumor and advanced stage ovarian cancer was confirmed. Patient had comorbidities like diabetes mellitus, hypertension and poor medical condition for a debulking surgery so we have decided to administer neoadjuvant chemotherapy. For that reason, due to the choice of senior surgeon, infracolic omentectomy, right salpingectomy, umbilical hernia repair was performed within the context of initial surgery. A weight and age adjusted dose of LMWH (4000 anti-Xa IU 40 mg enoxaparin sodium) was initiated for deep vein thrombosis prophylaxis during post-operative periods. Afterwards the surgery, chemotherapy treatment was planned according to pathology and patient was discharged with LMWH treatment; which as a protocol, was prescribed for a month period after surgery. Her final histopathology result was high grade serous ovarian carcinoma and thus she received chemotherapy (paclitaxel= 175 mg/m² and carboplatin= area under curve=6; per three weeks). After 4 cycles of chemotherapy, she underwent interval debulking surgery procedure, which was required and received the fifth chemotherapy cycle on 10th day of post-operative follow-up. Type 2 hysterectomy + left salpingo-oophorectomy + right oophorectomy + bilateral pelvic and paraaortic lymph node dissection + total omentectomy + appendectomy + splenectomy + tumoral tissue resection from diaphragm were performed within the context of interval surgical procedure. The following day of interval surgery, LMWH (4000 Anti-Xa IU/0.4 MI) was initiated for prophylaxis and the patient was discharged with the follow-up arrangements. On the 6th day of outpatient follow-up after receiving 5th cycle of chemotherapy, she was admitted to hospital with complains of nausea, vomiting, diarrhea and fever. Due to clinical examination as fever, oral aphthous ulcer was revealed; she was hospitalized for inpatient treatment and consulted to infectious disease department. The clinical status of the patient with normal hemato logic parameters was attributed to chemotherapy toxicity and gastrointestinal toxicity. IV antibiotics and fluid retention was administered and so clinical improvement achieved.

She has received the 6th cycle of chemotherapy and after 10 days of accomplished cycle, she underwent examination and laboratory results indicated that her platelet count declined to 9000/mm³ and the thrombocytopenia was revealed. Other parameters were in normal range. Thrombocyte replacement (6 units) was performed considering the hematological toxicity. After the replacement, platelet count has increased to 112.000/mm³. But the platelet count decreased to 34.000/mm³ within 6 days progressively. After that, thrombo cyt count slowly ascended to normal range within 15 days. She didn’t take LMWH yet in this period. She was accepted as chemotherapy-responsive based on the examinations she has undergone after 6th cycle of chemotherapy and therefore next planned 3rd chemotherapy cycle of the interval debulking surgery was decided not to be given.

She was hospitalized after 47 days of 6th cycle of chemotherapy due to infection. Leukocytosis was present in hematologic parameters with neutrophil dominance where platelet count was 118.000/mm³. The origin of the infection was thought to be the lymphocyte in abdomen thus IV antibiotics initiated. For prophylaxis due to hospitalization Clexane (4000 anti-Xa IU 40 mg enoxaparin sodium) was administered. Pulmonary embolism was developed on the 2nd day of inpatient follow-up so therefore heparin dose was increased to 2 x 6000 anti-Xa IU. After 5 days of LMWH use of treatment, a progressive decrease in platelets (>50%) was observed. Considering all of the findings, ‘HIT’ was suspected and hereby heparin treatment was stopped. Fondaparinux treatment initiated and laboratory tests applied. Platelet levels returned to normal limits within 6 days. Clinical and biochemical infection findings were resolved on the 7th day of IV antibiotics. Patient has taken fondaparinux treatment for more than 6 months.

Signed informed consent was obtained from the patient.

**Discussion**

Heparin-induced thrombocytopenia should be suspected in the patients who are currently receiving heparin or who received heparin and develops new onset of thrombocytopenia (i.e., platelet count <150,000/µL), a decrease in platelet count by 50 percent or more, even if the platelet count exceeds 150,000/µL, venous or arterial thrombosis, necrotic skin lesions at heparin injection sites and acute systemic reactions (e.g., fever/chills, tachycardia, hypertension, dyspnea, cardiopulmonary arrest) occurring after intravenous heparin administration (6,7).

Classically HIT related platelet decline is expected to occur within 5-10 days of heparin exposure but timing may show variability (8). Four types of clinical manifestations are defined in case of thrombocytopenia timing: 1. Typical-onset HIT, 2. Rapid-onset HIT, 3. Delayed-onset HIT and 4. Spontaneous HIT. Typical-onset HIT (70% of patients) manifests as a platelet count decrease that begins 5 to 10 days after starting a course of heparin. Rapid-onset HIT is the presenting feature in about one quarter of patients and is defined as a decrease in platelet count within 24 hours of administering a course of heparin. Delayed-onset HIT refers to when the platelet count begins to decrease after all heparin has been stopped. Spontaneous HIT is a rare disorder in which a patient develops a clinical profile suggestive of HIT and in whom high levels of HIT antibodies can be detected, even though no heparin has
been administered previously. The clinical progress of our case was mostly compatible with typical-onset HIT.

Factors such as; surgery, unfractionated heparin, female sex, older age increases the frequency of HIT. HIT occurs more commonly after exposure to UFH than exposure to LMWH in surgical patients (9). The pathophysiology of HIT may be different in unfractionated heparin than in LMWH because of the smaller molecule size of LMWH. (4) Ultra large complexes form most efficiently with UFH; their formation is 10-fold less efficient with LMWH and negligible with fondaparinux. These biochemical findings correlate with the greater incidence of HIT in patients receiving unfractionated compared with LMWH (1). This makes LMWH less likely to induce HIT than the larger unfractionated heparins (4). In a study of meta-analysis which aim was to determine the risk for HIT with unfractionated and low-molecular-weight heparin thromboprophylaxis, the absolute risk for HIT was only 0.2% with LMWH and was 2.6% with unfractionated heparin (9). Heparin dose is not relevant of the increasing risk of HIT. Therapeutic doses of heparin may result in a greater incidence of HIT than prophylactic doses, but there are no data that clearly define the relationship between heparin dose and clinical findings (10).

This case represents an example of HIT. In our case, progressive thrombocytopenia was observed after heparin treatment, pulmonary embolism was developed in consequence of thrombosis. Thus, HIT was suspected. The important point for clinicians about HIT is the decision to continue using heparin as a critical anticoagulant agent or cease the heparin therapy and choose an alternative non-heparin anticoagulant. The current treatment options for HIT are argatroban, fondaparinux, and bivalirudin. In the present case, fondaparinux was used.

A clinician should always consider clinical and laboratory evidence in evaluating patients for HIT. However, definitive laboratory data (i.e., immunoassay and/or functional assay for HIT antibodies) may not be available for several days. Since clinician needs to decide urgently, should make a preliminary diagnosis of HIT based on clinical findings and immediately available laboratory data. Clinicians can confirm or refute the diagnosis once they have the definitive results of HIT antibody testing. The 4Ts score may be used to have an early diagnose of HIT which the test includes; thrombocytopenia, the timing of the platelet decline, the presence of thrombosis, and other potential causes of thrombocytopenia (11) (Table 1). The system yields an integer score between 0 and 8 with scores of 0-3, 4-5, and 6-8 classified as low, intermediate, and high pretest probability for HIT, respectively (12). In a meta-analysis, the positive predictive value of high, intermediate, and combined high and intermediate probability 4Ts scores was 0.64, 0.14, and 0.22 respectively. The pooled estimates of sensitivity and specificity of the 4Ts at a cut-off of ≥4 were 0.99 and 0.54, respectively. The 4Ts score was 8 in presented case and accepted as high probability group.

In gynecologic oncology patients, there is no evidence data of HIT existence. However, in obstetrics patients which UFH was used for prophylaxis, HIT was seemed to have a 1.5% rate (13). Greer and Nelson-Piercy, in their studies, de-

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**Table 1: 4Ts Scoring System**

<table>
<thead>
<tr>
<th>4Ts Scoring Parameters</th>
<th>2 Points</th>
<th>1 Point</th>
<th>0 Point</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thrombocytopenia score</td>
<td>&gt;50% platelet decrease to nadir ≥20 x10⁹/L</td>
<td>30%-50% platelet count decrease or nadir 10-19 x10⁹/L</td>
<td>&lt;30% platelet decrease or nadir &lt;10 x10⁹/L</td>
</tr>
<tr>
<td>Timing of platelet count decrease, thrombosis, or other sequel (first day of heparin course =day 0)</td>
<td>Day 5-10 onset or ≤1 day (with recent heparin exposure within past 5-30 days)</td>
<td>Consistent with day 5-10 decrease, but not clear (e.g., missing platelet counts), or ≤1 day (heparin exposure within past 31-100 days), or platelet decrease after day 10</td>
<td>Platelet count decrease ≤4 days without recent heparin exposure</td>
</tr>
<tr>
<td>Thrombosis (including adrenal infarction) or other sequel (e.g., skin lesions)</td>
<td>Proven new thrombosis, or skin necrosis (at injection site), or post-IV heparin bolus anaphylactic reaction</td>
<td>Progressive or recurrent thrombosis, or erythematous skin lesions (at injection sites), or suspected thrombosis (not proven)</td>
<td>None</td>
</tr>
<tr>
<td>Other cause for Thrombocytopenia</td>
<td>No explanation for platelet count decrease is evident</td>
<td>Possible other cause is evident</td>
<td>Definite other cause is present</td>
</tr>
</tbody>
</table>

**Total score of current case**

| 8 | HIGH RISK |

The 4Ts score is the sum of the values for each of the 4 categories. Scores of 1-3 considered to correspond to a low, Scores of 4-5 considered to correspond to an intermediate, Scores of 6-8 considered to correspond to a high probability.
fined that none of the 2777 patients who were used LMWH as prophylaxis, HIT was not detected (14).

Individuals with suspected HIT should stop using heparin immediately and therefore patient should have a non-heparin treatment. Individuals with a confirmed diagnosis of HIT should continue the non-heparin anticoagulant approximately 3-6 months. The diagnosis of HIT is based on clinical features supported by laboratory testing which are immunoassays and functional assays. This case report describes a patient whom was suspected for the diagnoses of HIT because of the clinical and laboratory findings and therefore heparin treatment was cancelled immediately and another anticoagulant treatment was initiated. Importantly, one should not wait for thrombosis to develop before suspecting HIT, because thrombocytopenia often precedes thrombosis. Early intervention has the potential to prevent thrombotic events, which are the major cause of morbidity and mortality in patients with HIT.

Conclusion

HIT is a temporary condition and the platelet count returns to normal within a few days to several weeks. HIT may both occur by the use of LMWH or UFH for the prophylaxis of thrombosis. By the use of fondaparinux, the platelet levels come to normal within 2-9 days. Misdiagnosis of HIT can have adverse outcomes. When the suspicion of HIT emerges, the clinician should use clinical probability scores and confirm the diagnosis with a peripheral blood sample and laboratory tests. HIT emerges rarely but the treatment is difficult so the treatment should be approached with care.

Conflict of Interest Statement
The authors declare no conflict of interest

References


