Profound Thrombocytopenia after Treatment with Tirofiban; On-Pump Surgery for Acute Aortic Dissection and Coronary Bypass Surgery

Beckers Stefan1*, Remory Isabel1, Van Der Meeren Sam1, Bougie Daniel3, Demanet C2, Jochmans K2 and Poelaert Jan1

1Department of Anesthesiology and Perioperative Medicine, Universitair Ziekenhuis, Brussel, Belgium
2Department of Hematology, Universitair Ziekenhuis Brussel, Belgium
3Blood Center of Southeastern Wisconsin, Milwaukee, WI, USA

Abstract

We describe the case of a 69-year-old man who developed glycoprotein IIb/IIIa receptor (GPIIb/IIIa) antagonist associated thrombocytopenia and the successful management during urgent aortic dissection and coronary artery bypass grafting. A flow cytometric assay was developed to confirm the diagnosis of GPIIb/IIIa-antagonist induced thrombocytopenia. This case was further complicated by the concomitant presence of heparine/PF4 platelet antibodies, but was proven not to be the causal factor of the thrombocytopenia.

Keywords: Thrombocytopenia; IIb/IIIa glycoprotein inhibitors; Aortic dissection

Introduction

Problems related to the use of glycoprotein IIb/IIIa(GPIIb/IIIa) receptor antagonists, such as abciximab, eptifibatide and tirofiban, are not infrequent [1] and could have a major impact on the clinical course of an often critically ill patient. We describe a patient in whom severe thrombocytopenia was diagnosed, although the cause was initially unclear. Both heparin-induced thrombocytopenia and GPIIb/IIIa antagonist related antibodies obscured the picture, along with other possible causes of thrombocytopenia.

Case Description

A 69-year old male was transferred from a peripheral hospital to our University Hospital with suspicion of an interposterolateral infarction. He had a medical history involving arterial hypertension and a myocardial infarction, nineteen years ago. No allergy was known or any personal or familial bleeding history. Initially he was treated with acetylsalicylic acid, nadroparin, clopidogrel and tirofiban. The patient underwent a coronary angiography, revealing an occlusion of the left anterior descending and right descending posterior coronary artery. An aortic valve plasty was completed. The patient was weaned from cardiopulmonary bypass and hemodynamically stabilized with an intra-aortic balloon pump, dobutamine (5 microgram/kg.min) and norepinefrin (150 ng/kg.min). The postoperative course was complicated with the development of a right bronchopneumonia, which was treated with amoxicillin. The platelet count remained normal without need for further transfusion and the patient was dismissed from the hospital 14 days postoperatively.

Blood samples, taken before cardiopulmonary bypass, were analyzed to elucidate the cause of the acute thrombocytopenia. Flow cytometric detection of tirofiban-dependent antibodies was performed in our hospital [1]. In brief, platelet-rich plasma was isolated from acid citrate dextrose anticoagulated blood, procured from normal group O-donors. Platelets were further isolated by centrifugation in acid citrate dextrose anticoagulated blood, procured from normal group O-donors. Platelets were further isolated by centrifugation in

...
The patient was dismissed after 12 days of hospitalization. After the presence of heparin. The intervention occurred uneventful and angina and underwent coronary angiography and subsequent CABG.

A year later, the patient was again admitted with de novo (HIT), as described previously [2]. which permitted us to exclude heparin-induced thrombocytopenia as 0.5) while the functional flow cytometric test results were negative, which permitted us to exclude heparin-induced thrombocytopenia (HIT), as described previously [2].

After the intervention, the platelets number recovered quickly (Table 1). A year later, the patient was again admitted with de novo angina and underwent coronary angiography and subsequent CABG. Tirofiban was avoided and thrombocytopenia remained absent in the presence of heparin. The intervention occurred uneventful and the patient was dismissed after 12 days of hospitalization. After the procedure, the patient was informed and gave approval for the publication of the manuscript.

Discussion

GPIIb/IIa antagonists (abciximab, epftibatide, tirofiban) are widely used in the management of a variety of patients with acute coronary syndromes; they prevent platelet aggregation and thrombus formation, thereby improving outcomes of these patients [3]. Therapy with these agents may lead to bleeding complications and in rare cases to thrombocytopenia, challenging the perioperative management of these patients. In clinical trials with GPIIb/IIa inhibitors, mild to severe decline of platelet counts has already been shown [1,4,5]. In particular, this decline has been demonstrated for abciximab in ± 2%, often related to heterogeneous antibodies [6]. Tirofiban and eptifibatide show lower frequencies of associated thrombocytopenia [4,7].

This case demonstrates a profound acute thrombocytopenia after use of tirofiban, a GPIIb/IIa inhibitor. Antibodies against tirofiban were undiagnostically demonstrated in this patient. The differential diagnosis of the severe acute thrombocytopenia at initial presentation was pseudo-thrombocytopenia and two possible causes of drug-induced immune thrombocytopenia. Both heparin-induced thrombocytopenia (HIT) and tirofiban-related thrombocytopenia had to be excluded.

Pseudo-thrombocytopenia is defined as an in vitro clumping of platelets in blood samples anticoagulated with ethylenediaminetetraacetic acid (EDTA) [5,8]. In cases of pseudo-thrombocytopenia, there is neither in vivo thrombocytopenia nor enhanced aggregation. This artificial miscalculation of platelets was excluded after repeated low platelet counts in blood anticoagulated with citrate (Table 1) and because of the microscopical absence of platelet aggregates in blood smears. However, following Bizzaro [9], pseudo-thrombocytopenia can still occur in citrated blood samples. Ideally, to obviate this phenomenon, blood should be collected in ammonium oxalate and platelets counted in a Burker chamber.

HIT occurs due to the administration of the unfractionated or low molecular weight heparin and often leads to white thrombosis [10]. It typically presents at day 5-10 after first heparin administration, and the platelet count rarely falls below 20000/mm³ [11]. Because none of these criteria were met in our current case, the pre-test probability score for HIT was low.

However, cases of “fast-onset” HIT have been described and thus this potential cause had to be excluded [12]. IgG Heparin/PF4 antibodies were detected in the patient's serum, using an immunological HIT screening test (Zymutest HIA IgG, Hyphen Biomed). The activating potency of these antibodies was measured with a flow cytometric analysis.

Figure 1: Tirofiban-dependant antibody test; Flowcytometry result showing a shift in fluorescence intensity compared to platelets processed identically, except for the presence of drug. An aliquot was sent to the Blood Research Institute of Blood Centre of Wisconsin for confirmation.

<table>
<thead>
<tr>
<th>Time</th>
<th>20/11 10:30 A.M.</th>
<th>20/11 14:20 P.M.</th>
<th>20/11 15:30 P.M.</th>
<th>20/11 15:50 P.M.</th>
<th>20/11 04:30 P.M.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Platelet count 10³/mm³</td>
<td>265</td>
<td>3</td>
<td>3 (citr)</td>
<td>29 (citr)</td>
<td>30</td>
</tr>
<tr>
<td>PTT %</td>
<td>100</td>
<td>100</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>APTT sec</td>
<td>27</td>
<td>28.9</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Number of units of platelets</td>
<td>1</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hb g/dl</td>
<td>15.2</td>
<td>13.8</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Htc %</td>
<td>46</td>
<td>40.4</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 1: Perioperative laboratory values after tirofiban and platelets administration.
test, using CD62p (P-selectin) as a marker for platelet activation. This analysis was performed according to the method by Denys et al. [13]. Results were negative, thereby ruling out HIT. Moreover these antibodies can be excluded as cause of the thrombocytopenia from a clinical point of view: platelet levels restored to normal after withdrawal of tirofiban whereas the patient was kept on different drugs including heparin during further hospital stay.

Antibodies to heparin/PF4 complexes may exist in the absence of a clinical manifestation of HIT; in such case, the IgG heparin/PF4 antibodies present in the patient's serum are unable to activate the innate platelets. According to Warkentin and Greinacher, the variable platelet reactivity may correlate with the platelet Fc receptor-dependent reactivity and the existence of different polymorphisms of the receptor [10,13]. To elucidate this phenomenon, further effort was performed in our laboratory to identify the false positive results in the in vitro reaction or provide proof of the patient's platelet unresponsiveness.

The profound thrombocytopenia (3000/mm³) in our patient was provoked by the administration of tirofiban as shown by the flow cytometric detection of IgG solely in the presence of the drug tirofiban. The flow cytometry test (Figure 1) showed a shift in fluorescence in the presence of 4 µg/ml, indicating increased antibody binding in the presence of tirofiban.

No shift in fluorescence was observed in the absence of tirofiban. In this case, the patient's plasma was unable to activate a pool of responsive donor platelets in response to added heparin, excluding HIT as the cause of thrombocytopenia. Therefore, it seemed unnecessary to perform further tests with the patient's innate platelets.

This case demonstrates once more the difficult differential diagnosis of acute thrombocytopenia. It also underlines the need of caution and rapid detection of drug-dependent antibodies with respect to GPIIb/IIIa inhibitors, as these drugs will be used more frequently. The test is even more important in the light of the combined therapy with heparins, blurring the diagnosis in the presence of critical illness.

References