Management of Cholecystectomy in a Patient with Severe Von Willebrand Disease

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Abstract

Von Willebrand disease is the most common hereditary bleeding disorder; its Type 3 is the most severe form with a high risk for surgical bleeding, both intraoperatively and during the postoperative period. Despite that, there is a scarcity of data describing the management of these patients undergoing surgical procedures. We report on a 55-years-old woman with a long history of Von Willebrand disease type 3, who had an emergency laparoscopic cholecystectomy performed using VWF/FVIII concentrate (Wilate®), according to an adapted individual protocol, without significant bleeding or other complications. To our knowledge, this is the first case reported of a surgical procedure, under VWF/FVIII concentrate Wilate®, performed in a patient with VWD type 3 in Morocco.

Keywords

Von willebrand disease type 3, Von willebrand factor, factor VIII, VWF/FVIII concentrate Wilate®, Surgical bleeding

Introduction

Von Willebrand disease (VWD) is the most common hereditary bleeding disorder with a prevalence of approximately 1%-2%, affecting both genders equally [1]. It is a heterogeneous disease [2]; its Type 3 is defined by a total quantitative deficit in VWF. It is the most severe form of this disease, represents 5% of the forms of VWD [3] and is characterized by a low level of Von Willebrand Factor (VWF) and Factor VIII of coagulation (FVIII). The patients with VWD type 3 are at high risk for surgical bleeding, both intraoperatively and during the postoperative period. Despite that, there is a scarcity of data describing the management of patients with VWD type 3 undergoing surgical procedures. The treatment recommended in such situations is prophylactic admission of VWF/FVIII concentrate (Wilate®), according to an adapted individual protocol, without significant bleeding or other complications. To our knowledge, this is the first case reported of a surgical procedure, under VWF/FVIII concentrate Wilate®, performed in a patient with VWD type 3 in Morocco.

Case Report

A 55-year-old woman presented to the emergency department for a sudden onset pain in the right hypochondrium, radiating to the right shoulder, associated with nausea and vomiting. Her past medical history was remarkable with a long-standing of repeated epistaxis and menorrhagia, which were neglected by the patient. Until the age of 36 years old, the patient had presented a significant hemorrhagic syndrome following tooth extraction. This motivated a hematology consultation and the diagnosis of Willebrand disease type 3 was retained. From this age, the patient led a normal life under a follow up by hematology consultations in our institution. The physical examination located pain that blocked deep inspiration (positive murphy’s sign) and a localized defense in the right hypochondrium. An abdominal ultrasound was performed showing an enlargement of the vesicular wall with a vesicular lithiasis that was 1.2 cm long.

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IU/Kg or 2000 IU once a day for 10 days. The objective was to maintain adequately a VWF and plasma FVIII level above a threshold of 50% for at least 10 days. For lack of material resources, the patient only took 5 days of the maintenance treatment instead of the recommended 10 days (Figure 1). The hemostasis assessment, carried out after 48 hours of the loading dose, which is also the same day of surgery, showed a prothrombin time at 108%, Partial Thromboplastin Time Activated at 29 sec, the FVIII at 60%, vWF-Ag 66% and vWF-Rco at 24%. She recovered without any hemorrhagic incident, which is the reason tranexamic acid was not used. The patient was discharged from the hospital on the fifth days after the surgery. Considering the characteristics of her disease, we decided not to place the patient on anticoagulation therapy. Until

The results of coagulation tests are listed in (Table 1). They showed correct values of prothrombin time and fibrinogen. In addition, the platelet count was within normal range, but serum FVIII activity (FVIII: C) was abnormal, reduced to a 9% (normal range: 70-140). Von Willebrand factor antigenemia (vWF-Ag) and activity (vWF-Rco) were undetectable (normal range: 50-150%) with incalculable VWF: Rco/VWF: Ag report. The diagnosis of acute lithiasis cholecystitis was accepted. Given the persistence and the accentuation of the pain despite an analgesic treatment by intravenous route, an emergency laparoscopic cholecystectomy was indicated. Indeed, the surgical treatment is the curative treatment of the acute lithiasis cholecystitis, since it allows the management, simultaneously, of the cause (lithiasis) and the consequence (infection). For our patient, the risk of surgical bleeding is higher, both intraoperatively and during the postoperative period. It was believed that surgery could be performed safely if adequate levels of factor VIII could be maintained.

The patient received VWF/ FVIII concentrate (Wilate®; Octapharma AG, Lachen, Switzerland) according to an adapted individual protocol developed on the basis of a multidisciplinary consultation between the hematologist, the surgeon and the anesthetist-resuscitator, while being based on the recommendations published by the High Authority for Health [8].

The loading dose was 60 IU/kg or 4500 IU (for a weight of 70 kg) twice a day as a bolus, intravenously for two days. The maintenance dose was set at half the loading dose: 30 IU/Kg or 2000 IU once a day for 10 days. The objective was to maintain adequately a VWF and plasma FVIII level above a threshold of 50% for at least 10 days. For lack of material resources, the patient only took 5 days of the maintenance treatment instead of the recommended 10 days (Figure 1). The hemostasis assessment, carried out after 48 hours of the loading dose, which is also the same day of surgery, showed a prothrombin time at 108%, Partial Thromboplastin Time Activated at 29 sec, the FVIII at 60%, vWF-Ag 66% and vWF-Rco at 24%. She recovered without any hemorrhagic incident, which is the reason tranexamic acid was not used. The patient was discharged from the hospital on the fifth days after the surgery. Considering the characteristics of her disease, we decided not to place the patient on anticoagulation therapy. Until

| Coagulation tests                  | Patient values | Normal values |
|-----------------------------------|----------------|---------------|
| Clotting time (Sec)               | 11             | 10-13         |
| Prothrombin time (%)              | 140            | 70-140        |
| Partial Thromboplastin Time Activated (Sec) | 48.9     | 23-33         |
| Fibrinogen (g/l)                  | 2.75           | 2-4           |
| Facteur VIII activity (%)         | 9              | 70-140        |
| vWF: Ag (%)                       | Undetectable   | 50-150        |
| vWF: Rco (%)                      | Undetectable   | 50-150        |

Figure 1: Representativerecovery profile in our patient receiving multiple doses of wilate®.
now, the patient is always followed in consultations without showing hemorrhagic complications in connection to the surgical operation carried out.

Discussion

Von Willebrand disease is the most common bleeding disorder, but it is still unclear how to treat patients who undergo surgery to prevent or control the bleeding [4] since the risk of bleeding differs from one patient to another. Its evaluation is based on the patient’s own hemorrhagic risk, his comorbidities, the treatments that can modify hemostasis, and on the hemorrhagic risk of the procedure envisaged (surgical approach, infectious context …). This evaluation is the subject of a multidisciplinary consultation between the hematologist, the surgeon who will perform the surgery and the anesthesiologist-resuscitator, in order to write an appropriate individual therapeutic protocol.

Several countries have published recommendations: the United Kingdom [9,10], the United States [11], Italy [12] and the Netherlands [13]. All the recommendations that were made are based on observational studies or on a series of case studies. No randomized study and/or meta-analysis is available specifically concerning VWD. In addition, the classification of invasive procedures as “invasive procedure with low hemorrhagic risk” or “invasive procedure with high hemorrhagic risk” is debated in the literature. However, in general, we distinguish three main groups: Major (operations considered dangerous or life threatening such as orthopedic, general, we distinguish three maingroups: Major (operations considered dangerous or life threatening such as orthopedic, urological, urogynecology, and abdominal surgeries); minor (invasive procedures not considered life threatening, such as radioactively synovectomy and endoscopy); and dental procedures [4,6].

The use of factor concentrate is recommended for type 3. However, the important consideration in the use of this factor is the ratio of vWF: Rcof / FVIII: C in the concentrate, which differs in different commercial preparations. In this case, Wilate® (Octapharma AG, Lachen, Switzerland) was used exclusively in a patient how presents a VWD type 3 and underwent a laparoscopic cholecystectomy, which considered as a surgery at major risk of hemorrhage. It is a plasma-derived, albumin free, high purity concentrate of freeze-dried VWF and FVIII developed specifically for the treatment of VWD [7]. The manufacturing process preserves the integrity of the VWF triplet structure, and it contains two independent virus inactivation/removal steps with different modes of action. The VWF: Rco/fVIII: C ratio is close to the physiological value of 1:1, which should facilitate dosing and monitoring without any risk of accumulation of FVIII or thrombotic complications. This result was also reported in the study of Windyga, et al. [6] and indicates that there was no accumulation of FVIII with Wilate®. This is an important observation, because elevated plasmaFVIII levels have been reported to be associated with an increased risk of venous thrombosis and the risk is heightened in patients undergoing surgery [17,18]. No thromboembolic events were recorded in our patient or in a study of on-demand and prophylactic use of Wilate® [14].

Conclusion

The good control of hemostasis and close perioperative follow-up under VWF/FVIII concentrate are important to perform a major surgical procedure, safely, in vWD type 3 patients. To the best of our knowledge, this is the first case reported of a surgical procedure under VWF/FVIII concentrate Wilate®, performed in a patient with VWD type 3 in Morocco. This case provides encouragement for the performance of major surgical procedures with the support of VWF/FVIII concentrate even in severe cases of Willebrand diseases exposing the patient to a high risk of postoperative bleeding.

Conflicts of interest

There is no conflict of interest.

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