Disseminated intravascular coagulation developed after suction curettage in an adenomyosis patient: a case report and literature review

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Disseminated intravascular coagulation (DIC) is a high mortality coagulopathy. The basic diseases that can induce DIC include: obstetric events, metastatic malignancy, massive trauma and bacterial sepsis. But in this article, we reported on an extremely rare case of DIC in an adenomyosis patient who had undergone suction curettage on account of unwanted pregnancy. The patient had been successfully treated with transfusion of fresh frozen plasma and subcutaneous injection of nadroparin calcium instead of hysterectomy. A decrease in progestrone, bleeding in the adenomyosis lesions, tissue necrosis, releasing of tissue factors, activating of the exogenous coagulation system might play the crucial roles in the pathophysiology of acute DIC. Since early detection and treatment are essential for the prognosis of DIC, it is suggested to strengthen the monitoring of coagulation function in such patients.

Keywords
DIC, Early pregnancy, Suction curettage, Adenomyosis

1. Introduction

Disseminated intravascular coagulation (DIC) is induced by several clinical conditions such as infectious diseases and malignant diseases which account for about 2/3 of DIC. Obstetric events and trauma are also the main causes of DIC [1]. But DIC rarely occurs in patients with benign gynecological diseases. Adenomyosis is a chronic estrogen-dependent gynecological disease characterized by the presence of endometrial glands and stroma outside the uterine cavity [2]. In the present article, we reported a rare case of acute DIC after suction curettage due to unwanted early pregnancy in an adenomyosis patient. Although hysterectomy has been advised for women with severe symptoms from adenomyosis and also be a positive and effective treatment for acute DIC, there are still many patients with fertility requirements who have a strong desire of preservation of uterus. Approximately 20% of cases of adenomyosis involve women younger than 40 years [3, 4]. In the meantime, women are getting married later [5]. More and more patients with adenomyosis need to preserve their uteruses. In the present case, we provided a possibility that anticoagulant treatment could be considered as one option before considering of hysterectomy in such patients. The study was approved by Ethics Committee of Women’s hospital, School of Medicine, Zhejiang University (Approval No.: IRB-20200220-R).

2. Case report

The adenomyosis patient (37-year-old, gravidiy 4, parity 2) came to the emergency department of our hospital for severe abdominal pain and vaginal bleeding on October 19, 2018. The patient had suffered from adenomyosis for 5 years, but was not treated yet. Her menstrual flow and dysmenorrhea were tolerated. On October 15, 2018, she underwent suction curettage to terminate unwanted early pregnancy (8 weeks of pregnancy) in outpatient operation room of our hospital. There weren’t any drugs used during the suction curettage. But as a cervical preparation, 200 µg sublingual misoprostol had been taken by the patient 30 minutes before operation. The operation was successful. Approximately 10 mL blood was lost at the time of the suction curettage. The patient left the hospital after observation for 1 hour and there was little vaginal bleeding on the day of surgery. In addition to taking cefuroxime sodium 0.25 twice a day, no other drugs were taken after the operation. Then on the second day after operation, vaginal bleeding increased and abdominal pain intensified gradually. On the fourth day after operation, abdominal pain was unbearable and the patient came to our hospital for emergency treatment. In 2014 and 2016, she underwent cesarean sections. She had no medication history or family history associated with thrombotic diseases, and had not experienced any symptoms of bleeding tendencies.

At the time of consultation, the blood pressure and heart rate of the patient were in normal range. The body temperature was 37.9 °C. Physical examination showed pale skin and palpebral conjunctiva which indicated anemia. Pelvic examination revealed the uterus was as large as 3 months of gestation. And the uterus tenderness was obvious.

The results of a peripheral blood cell analysis were: white blood cell (WBC) count 25.7 × 10⁹ cells/L, neutrophil (N) percentage 93.6%, hemoglobin (Hb) 88 g/L and platelet (PLT) count 177 × 10⁹ cells/L. Coagulation test results were: prothrombin time (PT) 28.6 s, activated partial thromboplastin time (APTT) 69.4 s, thrombin time 44.6 s, interna-
ational normalized ratio (INR) 2.80, fibrinogen (Fg) < 0.6g/L. Serum biochemistry test results were: alanine aminotransferase (ALT) 34 IU/L, aspartate aminotransferase (AST) 38 IU/L, total bilirubin (TB) 37.8 μmol/L, direct bilirubin (DB) 10.1 μmol/L, indirect bilirubin (IDB) 27.7 μmol/L, creatinine 55.7 μmol/L. Trans-vaginal Ultrasound found that the size of the enlarged uterus was 8.1 × 7.2 × 7.3 cm and the posterior wall of the uterus was uneven and thickened.

We reviewed the preoperative laboratory test results of the patient and found no abnormalities in blood routine, coagulation function and serum biochemical tests. In order to eliminate the laboratory test error, we retested the peripheral blood cell analysis and coagulation function of the patient. And performed procalcitonin (PCT) test was done to exclude bacterial infection. The results were: WBC count 25.0 × 10^9 cells/L, N percentage 93.8%, Hb 76 g/L, PLT count 159 × 10^9 cells/L, PT 23.5 s, APTT 51.7 s, thrombin time 27.1 s, INR 2.18, Fb 0.6 g/L, D-Dimer > 20 mg/L, and PCT 0.09 ng/mL.

According to the diagnostic criteria of the ISTH (International Society on Thrombosis and Haemostasis) guidance [6] and the consensus of Chinese experts [7], DIC was diagnosed. Treatments including infusion of Tienam (Imipenem and Cilastatin sodium), transfusion of fresh frozen plasma, subcutaneous injection of nadroparin calcium were taken (Since the weight of the patient was 60 kilograms, the administered nadroparin calcium dose was 4100 IU every 12 hours for 3 days).

Meanwhile, abdominal CT and vascular ultrasound of extremities were performed. No obvious thromboses were found. However, CT showed an irregular low-density lesion with a range of 6.5 × 6.4 × 6.8 cm in the muscular layer of the posterior wall of the uterus. The finding suggested that the lesion of the posterior wall was necrotic cyst (Fig. 1).

Although the patient’s temperature was slightly high, according to the result of PCT, bacterial infection was excluded. There were no manifestations and evidences of malignant tumor, trauma or abnormal liver function. And in the light of common thrombosis area and symptoms of the patient, abdominal CT and vascular ultrasound of extremities were performed to detect mesenteric thrombosis and arteriovenous thrombosis of extremities. Based on a series of examinations and laboratory test results, the common causes of DIC were excluded. We considered that the cause of DIC in this patient was related to adenomyosis.

After treatment, the coagulation function of the patient began to improve 1 day later. Vaginal bleeding decreased. Abdominal pain relieved. Body temperature returned to normal. And the blood tests returned to normal range (Table 1). The patient was discharged 5 days later.

3. Discussion

Disseminated intravascular coagulation (DIC) is not an independent disease, but the ultimate common pathway of co-
agulation dysfunctions in many diseases. It is a clinicopathological syndrome. For the blood coagulation mechanism is diffusely activated, it promotes extensive fibrinogen deposition in small vessels, which leads to the damage of tissues and organs. On the other hand, the consumption of coagulation factors leads to the tendency of systemic hemorrhage. The two contradictions exist simultaneously in the development of DIC disease and constitute a specific clinical manifestation. Multiple organ dysfunction syndromes are the leading cause of death in patients with DIC. The basic diseases that can induce DIC include: systemic infection/severe infection, trauma, organ damage, malignant tumors, obstetric events, liver failure, etc. [6].

For patients after termination of pregnancy, sepsis caused by incomplete abortion is an important inducing factor of DIC. However, in the present case, transvaginal ultrasound founding ruled out the possibility of retained products of conception, PCT examination result ruled out bacterial infection. So in the present case, common causes of DIC, like serious infections, tumors, organ damage and liver failure, have been excluded based on a series of examinations and laboratory test results. The cause of DIC was considered relating to hemorrhage and necrosis of adenomyosis lesion after suction curettage.

Adenomyosis is a diffuse or localized lesion in which endometrial glands and stroma invade the myometrium of the uterus, presenting as hypermenorrhea and progressive dysmenorrhea [9]. It is commonly seen in women of reproductive age (especially in women from 25 to 45 years old). Only 2 reports (Contain 7 patients) of DIC associated with adenomyosis have been reported through our search for literatures in PubMed and Embase [10–16] (Table 2).

In above, 5 cases [10–13, 15] were menstruation-related. The other 2 cases [14, 16] were developed after dilation and curettage due to abortion. In the two cases, one patient with missed abortion underwent hysterectomy, the rest one in which the uterus was preserved was early pregnancy and similar to the present one we reported.

A typical sequence of intravascular coagulation and fibrinolysis includes the following: procoagulant exposure (such as tissue factor, TF), coagulation, fibrinolysis, end organ damage. Studies have shown that adenomyosis affects both the coagulation and fibrinolysis systems and the volume of adenomotic tissue may be associated with variations in the coagulation and fibrinolysis systems [17]. And tissue factors (TF) are highly expressed in adenomyosis [18]. Since TF is a procoagulant factor. We speculated that the following was the mechanism of the occurrence of DIC in this patient. A decrease in progesterone after abortion led to bleeding and necrosis in the adenomyosis lesions. Necrotic tissue of adenomyosis released a large number of TFs into the blood, which activated the exogenous coagulation system-microthrombosis. Blood was immersed in the muscular layer of the uterus which resulted in the separation, breakage and even degeneration of the muscle fibers, meanwhile, caused subsequent fibrin formation and microthrombosis inside adenomyosis tissues. Fibrin deposited on endothelial cells lining the walls of blood vessels and promoted infarct formation and thrombosis in peripheral vessels [17]. Microthrombosis that formed inside adenomyotic tissues lead to subsequent activation of the fibrinolysis system.

DIC is a dynamic process of ongoing thrombin generation and fibrinolysis. It can present as an acute, life-threatening condition or a chronic, subclinical process, depending on the degree and tempo of the process and the contribution of morbidities from the underlying cause. And a major principle in the management of DIC is treatment of the underlying cause in order to eliminate the stimulus for ongoing coagulation and thrombosis. As the menstrual related acute DIC case reported by Nakamura et al. [10], the underlying cause was adenomyosis, but the stimulus was menstruation. For the patient did not enter the stage of severe hyperfibrinolysis and terminal organ damage, when the stimulus was eliminated, surgical removal of the uterus was not required as anticoagulation therapy and supplementation of coagulation factors improved the condition. The present case showed an acute

| Laboratory data | 15/10 | 19/10 4:50 am | 19/10 5:46 am | 19/10 10:52 am | 19/10 21:51 pm | 20/10 | 21/10 | 22/10 | 24/10 |
|-----------------|-------|-------------|-------------|-------------|-------------|------|------|------|------|
| WBC (10^3/L)    | 9.1   | 25.7        | /           | 25          | 29.3        | 29.8 | 16.7 | 6.4  | 4.0  |
| N (%)           | 81.2  | 93.6        | /           | 93.8        | 97.8        | 96.1 | 83.6 | 68.3 | 58.6 |
| Hb (g/L)        | 118   | 88.0        | /           | 76          | 74          | 71   | 66   | 69   | 72   |
| PLT (10^9/L)    | 248   | 177         | /           | 159         | 138         | 146  | 176  | 198  | 294  |
| PT (s)          | 14.7  | 28.6        | 27.7        | 23.5        | 18          | 18.3 | 14.7 | 13.3 | 13.1 |
| APTT (s)        | 38.5  | 69.4        | 62.7        | 51.7        | 51.9        | 40.3 | 36.2 | 37.7 | 36.5 |
| Fg (g/L)        | 3.09  | < 0.6       | < 0.6       | 0.6         | 2.07        | 3.16 | 3.22 | 3.72 | 3.05 |
| INR             | 1.2   | 2.8         | 2.69        | 2.8         | 1.56        | 1.59 | 1.2  | 1.06 | 1.04 |
| D-dimer (mg/L)  | /     | /           | > 20        | > 20        | > 20        | > 20 | > 20 | 5.96 | 2.83 | 2.11 |
| ALT (IU/L)      | 32    | 34          | /           | 34          | /           | 31   | /    | /    | /    |
| AST (IU/L)      | 25    | 38          | /           | 38          | /           | 24   | /    | /    | /    |
| TB (µmol/L)     | 6.5   | 37.8        | /           | 37.8        | /           | 7.4  | /    | /    | /    |
| DB (µmol/L)     | 3.4   | 10.1        | /           | 10.1        | /           | 2.7  | /    | /    | /    |
| IDB (µmol/L)    | 3.1   | 27.7        | /           | 27.7        | /           | 4.7  | /    | /    | /    |

Results of peripheral blood cell analysis, coagulation function and liver function before and after termination of early pregnancy.
| Study            | Age | Onset period | Incipient symptoms                  | Laboratory data                                                                 | Preliminary treatment                      | Middle Outcomes                                                                 | Further treatment |
|------------------|-----|--------------|-------------------------------------|---------------------------------------------------------------------------------|--------------------------------------------|--------------------------------------------------------------------------------|--------------------|
| Nakamura Y 2002 | 32  | Menstruation | Dysmenorrhea; Gingival bleeding; Hematuria | WBC 18.2 × 10^9 cells/L, Hb 100 g/L, PLT 82 × 10^9 cells/L, PT 35.6 s, INR 7.28, APTT 94.2 s, fibrin/fibrinogen degradation products (FDP) 59.5 mg/dL, AST 46 IU/L, ALT 16 IU/L | Anticoagulation; Blood transfusion         | Laboratory values normalized. But prolonged APTT and PT was detected during the next menstrual cycle again. | GnRH agonist       |
| Jungmin S 2010  | 40  | Menstruation | Anuria; Elevated serum creatinine (Scr) level; Acute kidney injury | blood urea nitrogen (BUN) 41.1 g/dL, Scr 3.5 mg/dL, WBC 30.07 × 10^9 cells/L, Hb 100 g/L, PLT 39 × 10^9 cells/L, lactate dehydrogenase (LDH) 7914 IU/L, AST 329 IU/L, ALT 92 IU/L, TB 2.72 mg/dL, PT 24.6 s, INR 2.25, APTT 58.2 s, Fg 88 mg/dL, D-dimer 19.3 mg/L | Blood transfusion; Continuous renal replacement therapy; Therapeutic plasmapheresis Anticoagulation; | Blood transfusion; Laboratory values normalized. |
| Ohashi N 2011   | 51  | Menstruation | Nausea; Vomiting; Anemia; Thrombocytopenia; Acute renal failure | WBC 21 × 10^9 cells/L, Hb 108 g/L, PT 2.0 mg/dL, DB 0.3 mg/dL, AST 78 IU/L, ALT 20 IU/L, LDH 2317 IU/L, creatine phosphokinase (CPK) 181 IU/L, BUN 55 mg/dL, creatinine (Cr) 3.92 mg/dL, PT 16.2 s, APTT 30.1 s, Fg 410 mg/dL, D-dimer > 300 mg/L | Glucocorticoid; Plasma exchange; Hemodialysis | Lower abdominal pain and fever suddenly developed during the recovery course. Platelet count decreased dramatically. | Hysterectomy       |
| Yoo H J 2012    | 41  | Menstruation | Menometrorrhagia; Dyspea             | WBC 15.8 × 10^9 cells/L, Hb 88 g/L, Plt 52 × 10^9 cells/L, PT 23 s, INR 3.2, APTT 40 s, BUN 20 mg/dL, Cr 21.6 mg/dL, AST 52 IU/L, ALT 22 IU/L | Blood transfusion                          | Blood transfusion; Vaginal bleeding reduced. But the uterus enlarged gradually. | Hysterectomy       |
| Zhang J 2013    | 42  | After dilation and curettage | Lower abdominal pain; Continuous vaginal bleeding; Fever | WBC 12.92 × 10^9 cells/L, Hb 58 g/L, PT 19 × 10^9 cells/L, PT 13.2 s, APTT 34.4 s, Fg 200 mg/dL, FDP 694.9 mg/mL, D-dimer > 9999 mg/L, TB 51.4 mmol/L, DB 13.1 mmol/L, AST 126 IU/L, ALT 34 IU/L, LDH 2343 IU/L, BUN 14.36 mmol/L, Cr 276 mmol/L | Blood transfusion; Anti-infection           | Blood transfusion; Anti-infection           | Hysterectomy       |
| Cernogoraz A    | 56  | Menstruation | Abnormal uterine bleeding; Abdominal pain | WBC 20.01 × 10^9 cells/L, Hb 80 g/L, PLT 53.0 × 10^9 cells/L, APTT ratio 1.79, INR 1.28, Fg 145 mg/dL, Cr 2.24 mg/dL | Hemostasis; Blood transfusion; Anti-infection | Methrorragia ended. Coagulation improved. But creatinine elevated and anuria was ongoing. | Hysterectomy; Plasma exchange |
| Fuminori Kimura | 37  | After dilation and curettage | Abdominal pain; massive uterine bleeding; | WBC 19.2 × 10^9 cells/L, Hb 75 g/L, PLT 53.0 × 10^9 cells/L, C-reactive protein 24.1 mg/L, anti-thrombin-3 87%, thrombin antithrombin III complex 36.0 ng/mL, D dimer 111.1 μg/mL, plasminogen activator inhibitor-1 27.3 mg/mL, and plasmin α 2-plasmin inhibitor complex 13.7 μg/mL | Anticoagulation (nafamostat mesilate); Anti-infection | Abdominal pain and uterine bleeding improved rapidly. And the laboratory data normalized. | /                  |

* In Cernogoraz’s report, although hysterectomy was performed as soon as possible after transfusion therapy, the renal function was still not restored to normal levels 2 years after surgery. And specimen analysis revealed an enlarged uterus characterized by multiple intramural macroscopic thrombi. The microscopic view highlighted diffuse adenomyosis, as well as diffuse hemorrhage and the presence of microthrombi in smaller vessels.
Conflict of interest

The authors declare no conflict of interest.