**Abdominal Wall and Intra Pelvic Hematoma Presenting as Abdominal Pain after Short Course of Antibiotics in Patients on Long Term Warfarin Therapy**

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**Abstract**

Warfarin is the most commonly used anticoagulant in clinical practice. Despite its advantages it has serious side effects mainly bleeding due to narrow therapeutic range and several drug and food interactions. Spontaneous abdominal haemorrhage is one of the rare bleeding manifestations of it. Concomitant use of commonly used antibiotic are associated with an increased risk of bleeding among patients receiving warfarin.

We report two patients who presented with abdominal pain, later confirmed as abdominal wall and intra-abdominal hematoma by computed tomography of abdomen. Both of them were had concomitant use of antibiotics along with warfarin prior to this clinical presentation.

We highlight the importance of making a high degree of suspicion on abdominal haemorrhage in patients with warfarin treatment when presented as abdominal abdomen. We also emphasized the consideration of drug interaction of warfarin before prescribing other medication as well as close monitoring of INR during that period.

**Introduction**

Warfarin, the most commonly used anticoagulant worldwide, is indicated for the prevention of systemic embolism in patients with prosthetic heart valves, atrial fibrillation and for the primary and secondary prevention of venous thromboembolism [1,2].

Even though warfarin is efficient drug for the prevention of thrombo - embolism it is one of the most common drugs which have serious side effect mainly causing bleeding ranging from minor to fatal haemorrhagic manifestations [3-5]. Spontaneous abdominal wall and intra-abdominal haemorrhage is one of the rare bleeding manifestations of anticoagulant therapy.

All commonly used classes of antibiotics (Azole antifungals, Macrolides, Quinolones, Cotrimoxazole, Penicillin and Cephalosporin) are associated with an increased risk of bleeding among patients receiving warfarin [6,7].

We report two patients who presented with spontaneous abdominal wall and intra-abdominal hematoma with concomitant use of warfarin and antibiotics.

**Case History 01**

A 57 year old woman, on warfarin treatment following mitral valve replacement, presented with diffuse continuous mild to moderate abdominal pain and distension for 3-4 days. On the 4th day of symptom onset she noticed a dark pigmentation over the anterior abdominal wall.

She was able to maintain the INR in the therapeutic range most of the time during last 7 years. (Last INR level 1 month back was 2.6). She was prescribed Co amoxyclave 625 mg thrice daily, prednisolone 10 mg thrice daily and vitamin C 1 tablet thrice daily for respiratory tract infection 4 days prior to onset of symptoms.

On physical examination temperature 37.4°C, blood pressure 120/79 mm Hg, pulse rate 68 beats/min, and respiratory rate 12 breaths/min. A firm mass was palpable over the upper abdominal wall and it was not tender. She had ecchymosis in anterior abdominal wall (Figure 1). There was no evidence of free fluid on percussion.

Abdominal ultrasound revealed presence of multiple anterior abdominal wall masses and large intra pelvic hematoma measuring 10.9 cm × 9.8 cm × 9.8 cm adjacent to urinary bladder on right side. Computed tomography (CT) scan of abdomen confirms organized hematomas within the right rectus abdominis muscle in multiple levels (Figure 2) and a large hematoma in the pelvis (11.7 cm × 6.6 cm × 11 cm) to the right side of the urinary bladder displacing it (Figure 3).

Her other investigations on admission revealed INR level of 6.0, Hemoglobin level of 8.2 g/dL and serum creatinine of 1.53 μmol/L. Her platelet count, APTT, liver functions were within normal range.

Immediate suspension of oral warfarin treatment was done. She was transfused two pints of blood and pain managed with simple analgesia. Abdominal swelling resolved gradually in the following 4-5 days and no surgical intervention was needed. Her INR level came to therapeutic range 4 days after admission. Warfarin was restarted in smaller dose and adjusted according to the INR level. Her INR one

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week after admission was 2.6 and hemoglobin and serum creatinine levels achieved normal values.

**Case History 02**

A 58 year old female patient, with a 8-year history of dyslipidemia, essential hypertension and diabetes mellitus presented with localized anterior abdominal pain and swelling for 2 days duration. No history of trauma, urinary or bowel symptoms. Her long term medication includes atorvastatin 20 mg daily, losartan 25 mg bd, metformin 500 mg tds and warfarin 3.5 mg daily for last 5 years following prosthetic mitral valve replacement. She was prescribed co amoxyclav 625 mg tds, salbutamol 2 mg tds and piriton 4 mg nocte for respiratory tract infection 5 days prior to admission.

Physical examination was unremarkable except that moderate anemia and anterior abdominal wall lump measuring about 10 cm × 8 cm.

Laboratory investigation on admission showed INR level 10.0, Hemoglobin 6.4 g/dl with a hematocrit value of 61%. WBC count and platelet count, renal functions, liver functions and electrolytes were within normal range. Her abdominal ultrasound scan revealed hypo echogenic mass in anterior abdominal wall, which was confirmed as 8.1 cm × 4.3 cm ×12 cm (sagittal) size intra muscular hematoma within the right rectus abdominis muscle extending up to the pubic symphysis level in computed tomography (CT) scan (Figure 4).

She was managed conservatively and warfarin treatment withheld with daily monitoring of INR level. She was transfused 3 pints of blood during hospital stay.

Her INR level came to therapeutic range 4 days after stopping it. Warfarin treatment was reinitiated as small dose with adjustment of the dose during hospital stay. Her pre discharge INR level was 2.6 and hemoglobin level was 12.1 g/dL.

**Discussion**

Warfarin, the most commonly used anti-coagulant worldwide [1], is mediated through inhibition of vitamin K dependent gamma carboxylation of coagulation factors II, VII, IX, and X [8].

It is indicated for the prevention of systemic embolism in patients with prosthetic heart valves or atrial fibrillation and for the primary and secondary prevention of venous thromboembolism [2,9]. Despite its high clinical indication and efficacy serious side effects mainly causing bleeding led to its underutilization.

The most important factor causing the risk of bleeding is the intensity of anticoagulant therapy [5]. There are several factors including lot of drug and food interact the action of warfarin leading over anti coagulation, under anticoagulation or increased bleeding independent of changes in INR value [10,11]. Mechanisms associated with interaction include altered platelet action, alteration of vitamin K metabolism and alteration of warfarin metabolism. Commonly used antibiotics are well known to cause increased INR [6,7].

Although penicillins have not been shown to interact with warfarin [12], isolated few cases have been reported, particularly with broad-spectrum penicillins [13,14]. Our both patients were on Co-amoxyclyav for respiratory tract infection few days prior to development of abdominal bleeding.

Spontaneous abdominal wall hematomas are an uncommon cause of acute abdominal pain and are often misdiagnosed. It can also present as abdominal wall mass with or without ecchymosis [15,16].

CT scan can accurately confirm the diagnosis of abdominal wall and intra-abdominal hematoma [1].

The mainstay of management of abdominal wall hemorrhages is conservative including bed rest and analgesics. However, surgical treatment may indicate. In our patients the first measure was the
immediate suspension of oral anticoagulants with close monitoring of INR value and conservative management. None of them needed surgical treatment.

Conclusion

Keeping a high suspicion of abdominal haemorrhage is important if a patient on long term warfarin therapy presented with abdominal pain. Clinicians must consider the drug interaction of warfarin before prescribing other medication and need to monitor INR level closely during concomitant drug usage period.

References