



Bilateral Adrenal Hemorrhage in a Patient Anticoagulated with Apixaban for Antiphospholipid Syndrome (APS): A Rare Case Report

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Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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Case Report

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ABSTRACT

Context: Adrenal hemorrhage is a recognized yet rare complication associated with antiphospholipid syndrome (APS), an autoimmune disorder characterized by the presence of antiphospholipid antibodies that increase the risk of thrombosis. The incidence of adrenal hemorrhage in patients with APS has a diagnostic yield of approximately one-third, indicating that it may often go unrecognized in clinical practice. Traditionally, the management of APS has relied on anti-vitamin K therapy, such as warfarin. However, the advent of direct oral anticoagulants (DOACs) has prompted discussions about their use as an alternative. Recent clinical guidelines, however,

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strongly advise against the use of DOACs in patients with APS due to concerns over their safety and efficacy in preventing thromboembolic events.

Case Presentation: This case study describes a 53-year-old woman with a history of venous thromboembolism who was treated with apixaban, a direct oral anticoagulant. She presented with acute abdominal pain and symptoms suggestive of adrenal insufficiency. Initial evaluations revealed significant changes in her clinical condition, leading to further diagnostic imaging. A CT scan demonstrated bilateral adrenal hemorrhage, which was a surprising finding given her anticoagulation therapy.

Subsequent laboratory tests indicated a positive triple antibody profile, confirming a diagnosis of primary antiphospholipid syndrome. This case is particularly noteworthy as there have been limited documented instances of adrenal hemorrhage in patients receiving chronic anticoagulation with apixaban. The presentation of bilateral adrenal hemorrhage in the context of APS adds a crucial dimension to the understanding of the disease and its management.

Conclusion: This case highlights the potential for serious complications, such as adrenal hemorrhage, in patients with antiphospholipid syndrome, particularly those treated with direct oral anticoagulants. It underscores the necessity for clinicians to remain vigilant for such rare yet significant events in the management of APS. Additionally, this case raises important questions regarding the appropriateness of DOACs in this patient population, especially given the latest recommendations against their use. Increased awareness and further investigation into the implications of anticoagulation strategies in APS patients are warranted to improve patient outcomes and safety.

Keywords: Apixaban; antiphospholipid syndrome (APS); bilateral adrenal hemorrhage; case report.

1. INTRODUCTION

Antiphospholipid syndrome (APS) is a rare but serious condition.

The treatment of APS is based on the administration of anticoagulant medication, specifically anti-vitamin K (AVK), which has the ability to prevent the formation of blood clots.

The advent of direct oral anticoagulants (DOACs) in the fields of cardiology and vascular medicine has prompted questions regarding their efficacy in preventing thrombotic events in APS.

In light of the available literature, the latest French recommendations and the NASM advise against the use of DOACs in patients with APS.

Recent clinical guidelines strongly advise against the use of DOACs in patients with APS due to concerns over their safety and efficacy, especially in preventing thromboembolic events and avoiding complications like adrenal hemorrhage.

2. CASE PRESENTATION

The patient is a 53-year-old woman who is presenting with deep vein thrombosis of the left popliteal vein in relation to apixaban AOD (5 mg, administered in the morning and evening).

Following a one-month course of apixaban, the patient was admitted to the emergency department of Casablanca with respiratory distress and vomiting.

Upon admission, the patient was observed to be in a confused state. Her vital signs were notable for hypotension (99/70 mmHg), tachycardia (100 bpm), tachypnoea (24 bpm), and 90% free air saturation. Her Glasgow Coma Scale score was 14/15, and she exhibited no neurological deficits. Her temperature was 37.6°C. However, glucose levels were initially reported inaccurately at 0.29 g/L, without a proper reference to cortisol levels, which are crucial for diagnosis.

Following the administration of oxygen therapy via a high-concentration mask, correction of the hypoglycaemia and conditioning, the clinical examination revealed no additional heart murmurs and pleuropulmonary auscultation was normal.

A cardiac evaluation was conducted, which revealed no abnormalities in the electrocardiogram (Fig. 1).

The trans-thoracic echocardiogram (TTE) yielded borderline normal results, with a left ventricular ejection fraction (LVEF) of 60% (Fig. 2).

Biological assessment revealed the presence of hyponatremia at 129 mmol/L, hyperkalaemia at 5.9 mmol/L, and hypoglycaemia.

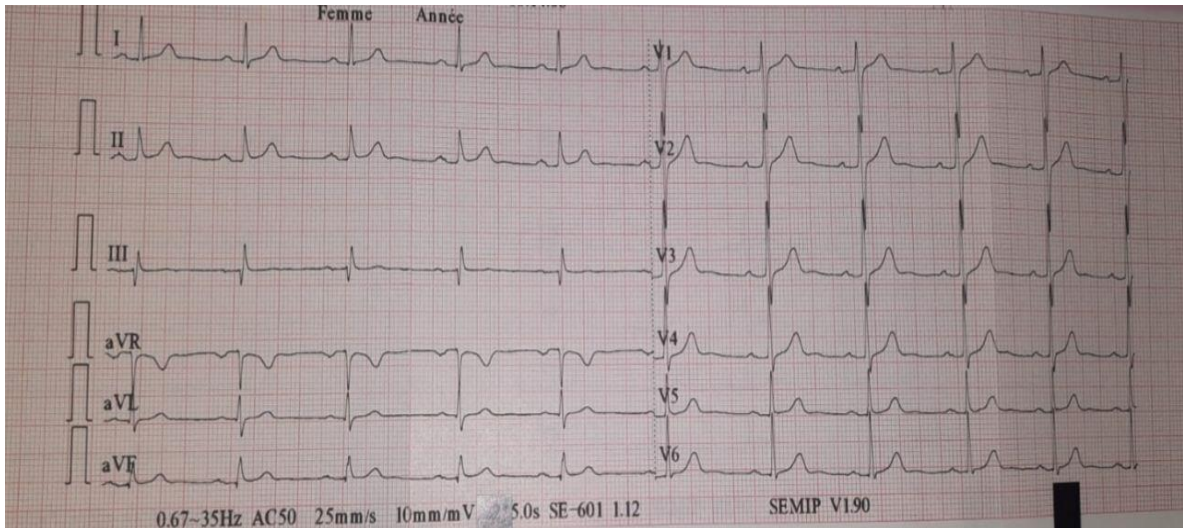


Fig. 1. The electrocardiogram (ECG) displayed sinus rhythm at a heart rate of 68 beats per minute (bpm), with no evidence of repolarisation abnormalities



Fig. 2. TTE revealed good biventricular function with an LVEF of 60%

In light of the clinical context, an abdominal CT scan was performed, which revealed bilateral haemorrhagic necrosis of both adrenal glands.

Management measures were initiated, comprising the administration of 100 mg of hydrocortisone hemisuccinate in conjunction with HDR.

The patient's clinical signs fully resolved, and her level of consciousness improved.

However, the lack of detailed discussion around adrenal insufficiency and cortisol measurements impairs the clarity of the case.

3. DISCUSSION

Adrenal haemorrhage represents a rare and potentially fatal complication of antiphospholipid syndrome (APS).

The mechanism for adrenal hemorrhage in APS patients is likely due to thrombosis within the

adrenal vasculature, leading to hemorrhagic necrosis [1].

The destruction of the adrenal cortex resulting from thrombosis and/or haemorrhage can occur rapidly; however, the extent to which it can affect adrenal function can vary.

The utilisation of a novel generation of direct-acting oral anticoagulants (DAAs) has largely superseded the use of warfarin in the management of hypercoagulability. DAAs possess predictable anticoagulant effects and do not necessitate dose monitoring, in contrast to warfarin. The convenience of DAAs has resulted in an increase in their utilisation. However, to date, there are no sufficiently robust randomised controlled trials that have established the safety of DAAs [1-8].

There is an ongoing debate about the use of DOACs in APS patients, with increasing evidence suggesting that DOACs may not be suitable for high-risk APS cases due to a higher incidence of thrombotic events. This case of adrenal hemorrhage further supports this notion, as it occurred while the patient was on apixaban.

The following factors have been identified as potential risk factors for HA: anticoagulant therapy, recent surgery, tumour metastases, corticotropin stimulation, adrenal tumour, physiological stress (trauma and burns), severe sepsis and APLS [2].

The selection of an anticoagulant for the treatment of antiphospholipid antibody syndrome (APS) may be pertinent to the risk of developing adrenal haemorrhage. However, the number of clinical cases available for analysis is limited, and therefore, it can only be hypothesised that warfarin may confer a lower risk of adrenal haemorrhage in comparison to other anticoagulant drugs (AODs).

Furthermore, it has been documented that patients with antiphospholipid antibody syndrome have experienced severe complications following a transition from warfarin to AOD therapy [3].

Adrenal haemorrhage has also been observed in patients without antiphospholipid antibody syndrome who were undergoing AOD therapy [4-5].

Currently, the evidence for secondary prevention of thrombotic events in patients with APLS using AODs is limited [6].

The European League Against Rheumatism (EULAR) has issued a recommendation against the use of AODs in APLS [7].

Until further evidence on the safety of AODs in APLS is available, warfarin should remain the reference anticoagulant [8].

4. CONCLUSION

APAS is a rare disease for which treatment is based on a consensus of experts.

The growing utilisation of AODs in a multitude of ailments gives rise to the question of their role in APS treatment. The available evidence suggests an increased risk of thrombosis in patients with high-risk APS, particularly in cases of triple positivity for antiphospholipid antibodies.

This has been demonstrated for rivaroxaban and a study is underway for apixaban (excluding arterial APS). Our case suggests that apixaban is ineffective in the treatment of high-risk APS, confirming the validity of recent recommendations. The discussion should be reorganized to start with a comprehensive explanation of APS, including its pathophysiology, the importance of preventing thrombotic events, and the pros and cons of available treatment options. DOACs, though effective in many other conditions, might not be suitable for APS patients, particularly those with high thrombotic risk.

Finally, this observation serves as a reminder that bilateral haemorrhagic necrosis of the adrenal glands is a serious complication of APS, which should always be investigated in the presence of abdominal pain with hypotension, in order to initiate specific treatment.

DISCLAIMER (ARTIFICIAL INTELLIGENCE)

Author(s) hereby declare that NO generative AI technologies such as Large Language Models (ChatGPT, COPILOT, etc) and text-to-image generators have been used during writing or editing of this manuscript.

ETHICAL APPROVAL

As per international standards or university standards written ethical approval has been collected and preserved by the author(s).

CONSENT

As per international standards or university standards, patient(s) written consent has been collected and preserved by the author(s).

COMPETING INTERESTS

Authors have declared that no competing interests exist.

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