Lenalidomide-associated arterial thrombosis in a patient with JAK2 positive atypical myeloproliferative neoplasm

Tanesh Kumar Ayyalu,1 Meltiady Issa2

DESCRIPTION
A 71-year-old woman, with a recent diagnosis of JAK2 positive atypical myeloproliferative neoplasm, presented to the emergency room (ER) for bilateral foot pain and purplish discoloration of her toes (figure 1). She started her first cancer treatment 3 weeks ago using lenalidomide with prednisone 20 mg daily. Venous thromboprophylaxis was not prescribed then due to anaemia and history of intracranial haemorrhage. ER work up included simple X-rays of her feet that were normal. Lower extremity ultrasound did not show any deep vein thrombosis, so she was discharged home. The patient did not have atrial fibrillation or hyperviscosity. No history of arterial thrombosis or smoking. Two weeks later, she was re-evaluated by her haematologist, and a hospital admission was arranged given worsening symptoms. Her physical exam now showed the purplish discoloration has progressed over both feet, associated with some toes turning black (figure 2). She had strong lower extremity pulses. Arterial lower extremity ultrasound showed normal ankle–brachial indices with no large vessel arterial disease. Heparin infusion was started given her gangrenous changes. Lenalidomide was discontinued. A CT angiogram of the aorta and large vessels looking for any embolic source was negative. Echocardiogram was normal. Her presentation was consistent with lenalidomide-induced arterial thrombosis. On discharge, she opted for a palliative approach with hospice. She passed away a month later.

Learning points
► While usually well tolerated, lenalidomide therapy can increase the thromboembolism risk, which, if not recognised early, can lead to significant complications.
► Thromboprophylaxis should be considered to prevent both venous and arterial events with lenalidomide therapy, based on risk factors and immunomodulators.
be catastrophic. Prevention and early recognition is key to improve morbidity and mortality.

Contributors We (TKA and MI) certify that we have participated sufficiently in the work to take public responsibility for the content, including participation in the concept, design, analysis and writing of the manuscript. We both have made substantial contributions to the work reported in the manuscript.

Competing interests None declared.

Patient consent Obtained.

Provenance and peer review Not commissioned; externally peer reviewed.

REFERENCES