

Case of the Irretrievable Inferior Vena Cava Filter

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Case Report

With increasing use of Inferior Vena Cava (IVC) filters, more reports are emerging of complications including dislocation of the filter [1,2]. We report the case of a 16 year old lady, who developed extensive right ileo-femoral deep vein thrombosis (with extension into Inferior Vena Cava [IVC]) and pulmonary embolism in a setting of oral contraceptive use. During her work up following presentation she was noted to be heterozygous for Factor V Leiden. Following diagnosis she underwent successful pharmaco-mechanical thrombectomy and balloon angioplasty of the involved ileo-femoral venous segment and IVC with consequent resolution of pain and swelling of her right leg. At that same time she also underwent placement of a prophylactic IVC filter (suprarenal Bard G2 filter [Bard Peripheral Vascular, Inc; Tempe, AZ]). Completion venogram demonstrated recanalization of the IVC and the right ileo-femoral segment. The filter was also demonstrated to be in good position. Subsequently, multiple attempts at removal of this temporary filter three months post placement were unsuccessful. The approaches applied included tandem techniques via trans-jugular and trans-femoral approaches. This was likely due to the fact that the filter had taken a horizontal disposition (prior to retrieval attempts) just above the level of the renal veins with herniation of struts into the left renal vein. Such a malpositioned filter brought with it the risk of migration and possible caval occlusion (7%-10% per decade) [3]. Additionally there was also the risk of renal vein thrombosis secondary to herniation of the struts into the left renal vein. While the patient had initially been asymptomatic (for 6 months following thrombolysis),

she went on to develop persistent flank and back pain. In the setting of a symptomatic malpositioned IVC filter who had previously failed multiple percutaneous attempts at filter removal, it was decided to proceed with open retrieval of the filter (10 months post placement). This was done via a right subcostal incision with the patient in a left lateral decubitus position. After adequate exposure of the perirenal inferior vena cava, control both proximally and distally of the IVC was obtained. The renal veins were also isolated and controlled subsequent to which the IVC was entered via an oblique incision and the filter retrieved. Delivery of the filter required careful dissection as it involved peeling away endothelialized struts from the surface of the IVC and renal veins. Some of the struts had to be cut away with a wire cutter. Following delivery of the filter, the defect in the IVC was repaired using a bovine pericardial patch.

The patient recovered quickly from this operation and at this time, 2 months post procedure she is back to her normal routine with complete resolution of symptoms.

References

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