Idiopathic Pulmonary Vein Thrombosis: A Case Report and Review of the Literature

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Citation

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Abstract

Pulmonary vein thrombosis (PVT) is a rare but potentially serious condition. Known etiologies of PVT include intrapulmonary neoplasm, postoperative complications after lobectomy, lung transplantation, as a complication of radiofrequency ablation and hypercoagulable state. Here, we describe an extremely rare case of idiopathic pulmonary venous thrombosis (PVT) in addition to reviewing the previously reported cases.

INTRODUCTION

Pulmonary vein thrombosis (PVT) is a rare but potentially serious condition. Known etiologies of PVT include intrapulmonary neoplasm, postoperative complications after lobectomy, lung transplantation, as a complication of radiofrequency ablation and hypercoagulable state. Here, we describe an extremely rare case of idiopathic pulmonary venous thrombosis (PVT) in addition to reviewing the previously reported cases.

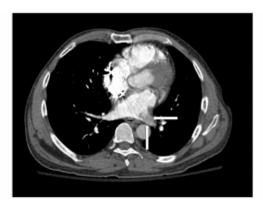
CASE REPORT

A 65-year-old male with past medical history of coronary artery disease, heart failure with preserved ejection fraction, symptomatic bradycardia status post permanent pacemaker, history of provoked deep venous thrombosis and pulmonary embolism (DVT/PE), seizure disorder and bilateral blindness due to glaucoma presented with syncopal episode. The patient was on his way to the bathroom when he started to feel sweaty and hot and then passed out. He denied any chest pain, palpitation, shortness of breath, nausea or abdominal pain. He denied any shaking movements, passing urine or tongue biting. His medications were aspirin, atorvastatin, levetiracetam, lisinopril and metoprolol. He has no family history of clotting disorders. Other than bilateral blindness, his physical exam was unremarkable including cardiovascular, neurological and respiratory exam. Blood work was unremarkable. EKG showed normal sinus rhythm. Echo was normal. Chest x-ray was within normal limits. Pacemaker interrogation showed no arrhythmias. A

subsequent CT pulmonary angiogram (CTA) (figure 1) revealed filling defect in the left lower pulmonary vein suspicious for thrombus. The patient was unable to have an MRI as he had a pacemaker. The patient was discharged on warfarin with a plan to repeat CTA and screen for thrombophilia, but the patient was lost to follow up.

Figure 1

The CT angiography reveals a well-defined filling defect in left lower pulmonary vein.



DISCUSSION

After reviewing the literature, we found only eight cases of idiopathic PVT have been reported, including the current case (see Table 1). Presenting symptoms in the series are not specific but mostly chest pain and shortness of breath.

Disease can occur in young and elderly. Males and females are equally affected. There are two reported complications related to idiopathic PVT: infarction of spleen and lung

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necrosis. CXR findings are not specific but may show infiltrate in lower lobes. Most of the cases diagnosed with using CTA. Other modalities used to confirm diagnoses are transesophageal echocardiography (TEE) and cardiac gated magnetic resonance imaging (MRI). Most of thrombus occur in right or left lower pulmonary veins compared to upper pulmonary veins. The appropriate treatment for PVT remains unclear but oral anti-coagulation appears to be the proper treatment. Duration of anticoagulation is unknown but among the eight cases, three had repeated CTA done. One of them showed resolution of thrombus after 3 months but the other two cases showed only partial resolution after 2 months.

CONCLUSION

Idiopathic PVT is rare disease with nonspecific presentation and potentially major complications. The best modality used for diagnosis is CTA. Most of the cases treated with oral anticoagulation but the duration remains unknown.

Table 1

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SeMildae SC:el.al. 2	32F	State and State	Acute left called abdowned pain, names and remiting	thinoun	operification in right lower listers	infection of spinor.	Contract enhanced abdustrial CT confirmed with ECG-gated 1403		Oral articogulation	Reguet CTA effor 2 months showed partial resolution
Amander et al.3	477	Unromorkable	Acate-theat pain and dyspress with Massive famosphysis	Unincurs	opecification of left lower lobe	Long necrosis	ET (high-resolution) acun confirmed intrasperative	Left referer pulmorany wite	Left lower lobe lobectomy. No long- term treatment	
Nomatos et al. 4	57 H	Opolipidemia	Classi pain with resourcible refunction	Negative	Not reported	Home	CT sheek, commeny anglogram. Se MDCT	Bilateral lower pulmonary within	Draf artitrospalation	
Hamil 1		coroney artery disease, coronery seturo logoes and compeditive head failure	scale shortness of breath	Dievated Isomocodoine	coedification of upper left-lobe	Name	CT angeography (CTN) shell	left superior palmonary With	Onal anticoaquilation (manifeste)	Repeat CTA after 5 months showed-complete resolution
We Fro		Unromorkable	chronic chest pain	Negative	urinoun	Home	CT anglography (CTII) chest	Left inferior palmoners sein entereling to airture	Left abid more resection and left lawer lobactoms plus long turns settings delice	
Ress.7	63 H	Unromarkable	substrate chest pain	Negative	to ecute changes	Home	CT angeography (CTN) cheet and Yes		One anticoogulation	
Tahu.0	55 M	CAD-with CABC and stant, CAG, symptometric lookpoints with pacemaker, accura- disorder	syncope	thinoun	ne acuto changes	Home	CT anglography (CTII) chest	left lower palmonery with	Onel entitrospolation (polation)	

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