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An Unusual Case of IJV Thrombosis

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Abstract: Internal jugular vein thrombosis is a rare but potentially life threatening condition occurring in the intracranial internal jugular vein or it's junction with the subclavian vein .This abstract presents a case study of a 50 year old obese patient with atypical chest pain and shortness of breath, diagnosed with internal jugular vein thrombosis. The investigation involved various tests ruling out ischemic causes and ultimately confirming subsegmental pulmonary thromboemboli. Thrombophilia work up was carried out and protein s deficiency was found. The patient responded well to anticoagulation therapy, and there is a need for further research to establish guidelines and standardised treatmentapproaches

Keywords: Internal jugular vein thrombosis, diagnosis, treatment, anticoagulation therapy, thrombophilia workup

1. Introduction

Internal jugular vein thrombosis refers to an intra luminal thrombus occurring anywhere from the intracranial internal jugular vein been to the junction of the internal jugular and subclavian vein to form the brachiocephalic vein. It may occur as a complication of and neck infections, surgery, central venous access, local malignancy, polycythemia, hyperhomocysteinemia, IV drug abuse

2. Case Presentation

50 year, old obese driver by profession was seen in cardiology outpatient department for atypical chest pain, shortness of breath with a subacute course 1 to 3 progressive over 3 weeks .He denies any history of smoking, recent surgery, trauma, infections, Intravenous drug abuse or catheter related procedures On examination he was found to have borderline high blood pressure (140/100 mm of hg, with BMI 32 ,had no systemic desaturation, non pulsatile JVP increase without any respiratory variation.

3. Investigations

Basic work up was done to rule out ischemic cause for chest pain by Electrocardiogram, 2D echocardiogram with no evidence of corpulmonale, Thread mill test, with minimal elevation of trop I 1.8ng/ml. Brain natriuretic peptide levels were normal with increased plasma dimers. So workup was done for Pulmonary thromboembolism. Computed tomography chest showing subsegmental Pulmonary thromboemboli. Neck vessel Doppler showing near total absent colour uptake on colour Doppler and echogenic foci filling the lumen of Right internal jugular vein which is decreased in caliber. Magnetic resonance venography was done to rule out intracranial extension of thrombus As there is no obvious risk factor for thromboembolic disease, we advised investigations with a thrombophilia work up Anti nuclear antibodies were negative, anti-nucleoproteinantibodies, antinucleosome antibodies, anticardiolipin antibodies, anti b2 glycoprotein were negative Antithrombin 3, homocysteine, protein c, factor 5 were within normal limits and were tested before introduction of anticoagulants, only proteins deficiency was found.

Treatment was initiated and in fractionated heparin for 48 hrs, symptoms regressed. Then patient discharged on oral novel anticoagulants after 5 days of admission. On follow up neck doppler showed partial recanalization and his plasma were negative at the end of 1month.

4. Discussion

Internal jugular vein thrombosis is a rare but potentially life-threatening condition that is often caused by central venous, catetherization, trauma, or infection. In this case, the patient had no risk factors for Internal jugular vein thrombosis. The management of Internal jugular veinthrombosis typically involves anticoagulation therapy with or without thrombolytic therapy.



CTPA image showing segmental & subsegmetal PTE

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5. Conclusion

Diagnosis and treatment with anticoagulation therapy can lead to successful resolution of the thrombus and prevent complications such as pulmonary embolism. The management of internal jugular vein thrombosis is heterogenous and currently based on the management of lower extremity deep-vein thrombosis, with a low rate of complications. The lack of guidelines and large series means that the modalities of diagnosis and treatment type and duration are variable. Given the low prevalence of internal jugular vein thrombosis, large, randomized studies would be hard to carry out. Data from our literature review suggest that treatments could tend towards those used for lower limb deep-vein thrombosis, but that the modalities of diagnosis and the duration of treatment and follow-up should be clarified

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