Case Report

Superior vena cava syndrome as a paraneoplastic manifestation of soft tissue sarcoma

Hosein Ghorbani, Mohsen Vakili Sadeghi *, Tahereh Hejazian, Majid Sharbatdaran

Babol University of Medical Sciences, Babol, Iran

Article Info

Article history:
Received 3 May 2017
Accepted 11 September 2017
Available online 17 February 2018

Introduction

Superior vena cava (SVC) syndrome occurs because of SVC obstruction; the obstruction can be due to external pressure on the vein or to an internal obstruction with thrombus formation in the vein. 1, 2 With the growing use of intravenous catheters and other devices, benign etiologies of SVC syndrome have become more common. 3, 4 Most etiologies of SVC syndrome are malignant with the most common being lung cancer, lymphomas (Hodgkin's and non-Hodgkin), and breast cancer. 1, 2 Other benign causes include thyroid goiter, aortic aneurysm, tuberculosis and thymoma. 2 In this article, we present the case of a woman with a malignant cause of SVC thrombosis but without any mass identified in the mediastinum.

Case report

A 46-year-old woman was admitted into hospital because of dyspnea, right arm pain and dysphagia. Forty-five days before admission she had developed pain in the right trunk and right scapula. Fifteen days prior to admission, she developed periorbital edema that slowly progressed to the entire face (Figure 1), neck, upper right extremity, upper trunk and both breasts. She also complained of erythema and pruritus of her forehead, cheeks and skin of the upper trunk during this period. She had no history of fever, weight loss or night sweating. In the physical examination, the patient was not febrile and her respiratory rate was 22 breaths per minute. Beside the edema, collateral vessels were visible in her upper chest.

A chest X-ray was normal. Color Doppler ultrasonography of the upper extremity veins revealed normal flow in the left jugular vein, non-obstructive echogenic thrombosis in the right jugular vein and obstructive thrombosis in the right subclavian vein. A spiral chest computed tomography scan with contrast enhancement revealed bilateral pleural effusion in particular on the right side; thrombosis was identified in the superior vena cava that extended to the left subclavian vein. Moreover, bilateral lung nodules were found compatible with metastasis, as was a mass in the right rotator cuff muscles with scapular bone destruction (Figure 2).

* Corresponding author at: Department of Hematology and Medical Oncology, Clinical Research Development Unit of Rouhani Hospital, Babol University of Medical Sciences, Babol, Iran.
E-mail address: mvakili89@gmail.com (M. Vakili Sadeghi).

https://doi.org/10.1016/j.htct.2017.09.001
2531-1379/© 2018 Associação Brasileira de Hematologia, Hemoterapia e Terapia Celular. Published by Elsevier Editora Ltda. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).
Laboratory data of the patient are shown in Table 1. Enoxaparin therapy was initiated and core needle biopsy of the mass was performed. The pathology exam identified sarcoma which was confirmed by immunohistochemistry staining (Figure 3). Fifteen days after starting chemotherapy, her upper trunk edema was reduced significantly but the patient suddenly developed dyspnea and expired before any diagnostic procedure was performed. Pulmonary embolism was the most probable cause in spite of the regular enoxaparin injections from the time of her admission.

Discussion

Intraluminal obstruction of the SVC may have benign or malignant causes. Long-term indwelling catheters, pacemaker wires and other intraluminal devices are common benign causes of SVC thrombosis.1–4

Because the contrast enhanced computed tomography scan of our patient did not identify any mass, it follows that the SVC syndrome was a paraneoplastic syndrome. Santra et al. reported paraneoplastic thrombus formation in SVC due to lung cancer.5 Moreover, paraneoplastic causes of SVC syndrome due to thrombosis have been reported in association with renal cell carcinoma,6 ovarian cancer7 and Richter
Obstruction of the SVC can also be due to intraluminal metastasis without thrombus formation. Takayoshi et al. reported a case of adenosquamous carcinoma of the duodenum with intraluminal SVC metastasis. Furthermore, invasive thymoma and prostate cancer with intravenous metastasis have been reported. Thus, when a cancer is diagnosed but without any mass being identified in the upper mediastinum, paraneoplastic thrombosis is the most probable cause of SVC syndrome as in our case unless anticoagulation was ineffective for which a biopsy of the intravenous lesion is necessary. In addition, when a cause for SVC syndrome is not found, a biopsy is necessary for differential diagnosis, which includes metastasis, thrombosis, granuloma, fungal lesion, etc.

Most common symptoms are shortness of breath, cough, and swelling of face, neck, upper chest and extremities. Swelling may result in stridor, dysphagia and hoarseness. Chronic SVC syndrome causes distention of collateral veins, which may be seen in the upper chest. Pleural effusion is observed in 60% of cases. The signs and symptoms of our patient were compatible with chronic SVC syndrome.

Management includes chemotherapy or radiotherapy for malignant causes when a mass is the cause. Stent placement is increasingly being used to ameliorate compression of the vessel resulting from malignant causes. Occasionally bypass surgery is performed. Other nonspecific methods include head elevation, mild diuresis and corticosteroids to decrease swelling and dyspnea. When thrombosis is the cause of SVC syndrome, thrombolysis and/or anticoagulation may be indicated. Our patient received anticoagulation and chemotherapy. Acute dyspnea and death were most probably due to pulmonary emboli. In the study of Paolo et al. of 842 patients with deep vein thrombosis, 181 were known to have cancer and recurrent thromboembolism occurred in 20.7%, most commonly during the first month of anticoagulation as in our patient.

**Conclusion**

We reported an uncommon cause of SVC syndrome due to paraneoplastic SVC thrombosis, with a poor outcome, most probably due to pulmonary embolism during anticoagulation. Physicians should be alert during the management of SVC syndrome, in particular when the cause is malignant.

---

**Figure 3** – Histology and immunohistochemistry of the tumor. (A) Wright’s and Giemsa staining. (B) CD34 staining. (C) Smooth muscle actin (SMA). (D) Desmin. (E) Ki-67 protein.
**Table 1 – Laboratory tests of the patient.**

<table>
<thead>
<tr>
<th>Variable</th>
<th>Value</th>
<th>Reference value</th>
<th>Variable</th>
<th>Value</th>
<th>Reference value</th>
</tr>
</thead>
<tbody>
<tr>
<td>WBC (/μL)</td>
<td>$8.1 \times 10^3$</td>
<td>$4.0 \times 10^3$</td>
<td>TSH (mIU/L)</td>
<td>2.8</td>
<td>0.34–4.25</td>
</tr>
<tr>
<td>Hemoglobin (g/dL)</td>
<td>13</td>
<td>12.0–15.8</td>
<td>PT (S)</td>
<td>15</td>
<td>12–14</td>
</tr>
<tr>
<td>Platelet (/L)</td>
<td>$306 \times 10^3$</td>
<td>$165–415 \times 10^3$</td>
<td>PTT (S)</td>
<td>27</td>
<td>25–35</td>
</tr>
<tr>
<td>Cr (mg/dL)</td>
<td>1.0</td>
<td>Female 0.5–0.9</td>
<td>Sodium (meq/L)</td>
<td>133</td>
<td>135–145</td>
</tr>
<tr>
<td>Calcium (mg/dL)</td>
<td>9.0</td>
<td>8.5–10.2</td>
<td>Potassium (meq/L)</td>
<td>4.0</td>
<td>3.5–5.5</td>
</tr>
<tr>
<td>Alp (U/L)</td>
<td>Normal</td>
<td></td>
<td>Blood sugar (mg/dL)</td>
<td>87</td>
<td>70–110</td>
</tr>
<tr>
<td>ESR (mm/h)</td>
<td>25</td>
<td>0–20</td>
<td>LDH (IU/L)</td>
<td>799</td>
<td>(Upper-limit = 480)</td>
</tr>
<tr>
<td>CRP (mg/L)</td>
<td>36</td>
<td>&lt;10</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

WBC, White blood cell count; Cr, Creatinine; Alp, Alkaline phosphatase; ESR, Erythrocyte sedimentation rate; CRP, C-reactive protein; TSH, Thyroid-stimulating hormone; PT, Prothrombin time; PTT, Partial thromboplastin time; LDH, Lactate dehydrogenase.

**Conflicts of interest**

The authors declare no conflicts of interest.

**Acknowledgement**

We thank the Clinical Research development unit of Rouhani Hospital of Babol.

**References**