Pulmonary Thromboembolism Followed by Deep Vein Thrombosis In A Young Man With G6PD Deficiency After ChAd0x1 nCoV-19 Vaccine Administration

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Case Report

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ABSTRACT

Recently the term vaccine-induced immune thrombotic thrombocytopenia (VITT) used for individual which have thrombotic phenomena followed by ChAdOx1 nCoV-19 vaccine (AstraZeneca) administration against SARS coronavirus. Here we report the 27 years old healthy male and known case of G6PD deficiency which come to emergency department with progressive right calf swallow from 12 days ago and hemoptysis from a day ago. He mentioned he had administrated First dose of AstraZeneca vaccine for 3 weeks ago. He admitted with suspected pulmonary thromboembolism (PTE) followed by Deep vein thrombosis (DVT). In color Doppler study there are dilation in right calf vain with elevated lab measurement d-dimer indicated DVT also in computed tomography angiography (CTA) there are some evidence of filling defect in left pulmonary branch and right inferior lobar artery which represent to PTE.

CASE DESCRIPTION

Here we present A 27-year-old young healthy man with a known case of G6PD deficiency, with pain in his right calf, from twelve days earlier and hemoptysis from one day earlier who was admitted in Taleghani Hospital in Abadan (Iran) for further work-up to rule out pulmonary thromboemboli and deep vein thrombosis. Symptoms had been initiated ten days after the administration of first dose of ChAdOx1 nCoV-19 vaccine (AstraZeneca). The pain was more severe at nights and involved whole part of the calf and became aggravated with exercise and activity. Hemoptysis was with flecks of blood in sputum. His past medical history indicated gastric bypass surgery in previous year as a treatment for Grade III fatty liver disease and obesity. He had no history of bleeding nor recent thrombotic accident.

In his social history, he mentioned smoking cannabis and cigarettes (one and half pack per day) since approximately five years ago. He was also using Tramadol tablet (50mg per day) since a year earlier. In his family history, his brothers also had cases of G6PD deficiency. The remarkable findings in his physical examination were tenderness in right calf and hemoptysis. In his first day of hospital admission, Heparin (1600 IU per hour) was initiated for suspected vein thrombosis (VT). Color doppler sonography for his right lower limb (Venous system) was performed to confirm the diagnosis which demonstrated dilation and loss of compressibility in superficial femoral vein (SJV; saphenous vein) and popliteal vein (PV), indicating extensive acute thrombophlebitis. Computed Tomography Angiography (CTA) indicated filling defect in left pulmonary artery with developing to inferior lobar branch and right inferior lobar artery which propounded PTE (Fig. 1). Three plus Qualitative Reactive Protein (CRP) measurement was also achieved. Peripheral Blood Smear (PBS) showed normal size, normochromic and normal morphology red blood cells without schistocytes, normal white blood cell count and morphology with left shift till band cell production stage.
Moreover, urine analysis showed slight hemoglobinuria. Second day Heparin was prescribed due to elevation in active Partial Thromboplastin Time (aPTT) and the patient started taking Rivaroxaban (15mg twice per day) and two units of Fresh Frozen Plasma (FFP) was ordered for him. Blood test revealed thrombocytopenia (79x103/mm3), elevated D-dimer test and marginally increase in alanine aminotransferase (Table 1). Echocardiography was also done for suspected right heart dysfunction, which the result was normal. Next day, a negative RT-PCR result rejected probable COVID-19.

**Table 1.** Laboratory results of a patient presenting with VITT; (ER; Emergency room/ HA; Hospital Admission/ PT; Prothrombin Time/ PTT; Partial thromboplastin time/ INR; International normalized ratio/ AST; Aspartate transaminase/ ALT; Alanine transaminase/ ALK-P; Alkaline phosphatase/ CRP; C- Reactive Protein/ CK-MB; Creatine Kinase-MB/ BUN; Blood urea nitrogen/ Cr; Creatinine/ LDH; Lactate dehydrogenase).

<table>
<thead>
<tr>
<th>Time line Lab data</th>
<th>ER admission</th>
<th>Day2 HA</th>
<th>Day3 HA</th>
<th>Day4 HA</th>
<th>Day5 HA</th>
<th>Day6 HA</th>
<th>Day7 HA</th>
<th>Day8 HA</th>
<th>Day9 HA</th>
<th>Day10 HA</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hemoglobin (Ref: 14-18) (g/dl)</td>
<td>13.1</td>
<td>11.8</td>
<td>11.3</td>
<td>11.7</td>
<td>11.8</td>
<td>11.8</td>
<td>12</td>
<td>11.7</td>
<td>12.1</td>
<td>12.5</td>
</tr>
<tr>
<td>Platelet count (Ref: 130-400) (×10^3/mm³)</td>
<td>101</td>
<td>79</td>
<td>141</td>
<td>137</td>
<td>141</td>
<td>140</td>
<td>153</td>
<td>164</td>
<td>186</td>
<td>166</td>
</tr>
<tr>
<td>Leukocyte (Ref: 4-10) (neut%) (×10³/µl)</td>
<td>7.6 (85%)</td>
<td>9.67 (59%)</td>
<td>7.7 (88%)</td>
<td>7.8 (57%)</td>
<td>8.38 (69.3)</td>
<td>9.3 (69%)</td>
<td>10.03 (67%)</td>
<td>11.67 (70.2)</td>
<td>8.99 (61%)</td>
<td>8.65 (61%)</td>
</tr>
<tr>
<td>D-dimer (Ref: ≤250) (ng/mL)</td>
<td>-</td>
<td>3327</td>
<td>-</td>
<td>3401</td>
<td>-</td>
<td>3412</td>
<td>-</td>
<td>3397</td>
<td>-</td>
<td>3456</td>
</tr>
<tr>
<td>PT (Ref: 11-13) (seconds)</td>
<td>18</td>
<td>25.6</td>
<td>16.8</td>
<td>17.7</td>
<td>17.6</td>
<td>21.2</td>
<td>14.8</td>
<td>21.5</td>
<td>19.3</td>
<td>21</td>
</tr>
<tr>
<td>INR (Ref: ≤1.1)</td>
<td>1.7</td>
<td>2.6</td>
<td>1.4</td>
<td>1.5</td>
<td>1.5</td>
<td>1.8</td>
<td>1.15</td>
<td>2.0</td>
<td>1.8</td>
<td>2.1</td>
</tr>
<tr>
<td>PTT (Ref: 25-45) (seconds)</td>
<td>90</td>
<td>≥120</td>
<td>42</td>
<td>44</td>
<td>45</td>
<td>36</td>
<td>35</td>
<td>56</td>
<td>60</td>
<td>56</td>
</tr>
</tbody>
</table>

Fig. 1. Filling defect is seen in the left pulmonary branch extending to the lower lobar branch and also in the right lower lobar branch extending to the sub-segmental branches.
The patient was hospitalized for ten days, Rivaroxaban (15mg BID) with initiation vitamin C were ordered for him. Symptoms were improved gradually while pain and swelling in the right calf were also reduced. In all those ten days, the patient was oriented and reliable. His vital signs were all in normal range with no dyspnea. The patient was discharged with Rivaroxaban (15mg twice a day), which changed to 20 mg daily after one month. As follow up, color doppler ultrasonography was performed which revealed normal popliteal vein and localized SFV partial thrombosis.

DISCUSSION

Today vaccination has become the main part of controlling COVID-19 pandemic [1]. According to recent studies, the administrations of COVID-19 vaccine produced by AstraZeneca have indicated some major thrombosis phenomenon [2]; most of them are grouped into sub massive thrombosis. In this case report, we presented a young healthy man without any risk factors of thrombosis and any past thrombosis event, who came in with massive bilateral PTE and parenchymal infarction in spiral computed tomography of the chest (Fig. 2).

Fig. 2. Parenchymal infarction in spiral computed tomography of the chest.
He was hospitalized for ten days, and Rivaroxaban tablet was prescribed for him. The Patient was discharged in good health condition and was followed up by a pulmonologist every month. Previously, thrombosis event was considered as a life-threatening event mostly for females, whereas it can occur in males too. This case was reported to document a potential serious side effect of AstraZeneca’s COVID-19 vaccine which is now an important parameter for controlling COVID-19 pandemic.

CONFLICT OF INTEREST

The authors declare that there is no conflict of interest.

REFERENCES