

Case Report
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Cervicofacial Herpes Zoster Complicating Ganglion-Pulmonary Tuberculosis: Case Report and Review of the Literature: A Case Report

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ABSTRACT

Tuberculosis is an infectious disease; it has a variable degree of presentation, most often pulmonary while the extrapulmonary location is dominated by lymph node involvement, especially cervical. On the other hand, herpes zoster results from the reactivation of the virus that causes varicella. Here, we report the case of cervicofacial herpes zoster complicating ganglion-pulmonary tuberculosis in a 19-year-old female patient with a history of varicella during childhood, who presents bilateral lymphadenopathy, the diagnosis of pulmonary and lymph node tuberculosis is confirmed by histology, she has treated with anti-tuberculosis drugs and the evolution was marked by the presence of lymphadenopathy and the appearance of cervicofacial herpes zoster.

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Introduction

Tuberculosis remains a major infectious scourge, particularly in developing countries. In 25% of cases, the location of tuberculosis is extra-pulmonary, the cervical lymph node involvement of which is the most frequent. Tuberculous lymphadenopathy is most often unilateral, single, and isolated. Cervical localization is by far the most frequent [1,2,3]. Lymph node tuberculosis is an advanced form of tuberculosis, always preceded by a primary pulmonary infection. The causative bacterium is called Mycobacterium tuberculosis or Koch's bacillus [1].

On the other hand, herpes zoster infection is a painful rash along a nerve or nerve ganglion. These rashes occur as a result of the reactivation of the virus that causes varicella; the varicella-zoster virus (VZV) most often affects the chest or face [4]. We report the case of ganglion-pulmonary tuberculosis complicated by cervicofacial herpes zoster.

Case Report

Four months ago, a 19-year-old patient with a history of normochromic normocytic anemia, varicella during childhood,

suffers from dyspnea stage 3; productive cough, weight loss, night sweats, right and left submandibular swelling. On clinical examination, we find lymphadenopathy of the territory 2 and 3 bilateral, mobile to the two planes, painless, and the largest of which on the left measures nearly 2 cm. The rest of the clinical examination was unremarkable.

Radiologically, the chest x-ray showing a left para-hilar opacity; Thoracic CT had demonstrated heterogeneous supra and subdiaphragmatic polyadenopathies with pulmonary parenchymal involvement and nodular splenomegaly, suggesting primarily granulomatous involvement with the cervical ultrasound showing bilateral polyadenopathy of territory 2 and 3. In addition, the search for BK in the sputum is positive.

Histologically, a cervicotomy was performed, suggesting an inflammatory granulomatous necrotizing reaction compatible with a tuberculous origin. The diagnosis of pulmonary and lymph node tuberculosis was retained. The anti-tuberculosis treatment was started according to the 2RHZE and 7RH protocol. The evolution was marked by the persistence of the right and left cervical lymphadenopathies; a cervical-thoracic-abdominopelvic CT (figure:1) for control showing supra and sub-diaphragmatic lymphadenopathy with hepatomegaly and nodular splenomegaly

forming part of a granulomatous involvement, cervical ultrasound shows hypoechogenic globular polyadenopathies including only one partially calcified of the right basi-cervical site, and the appearance of vesicular lesions (figure:2), suggesting primarily a right cervicofacial zoster on the same side lymphadenopathy.

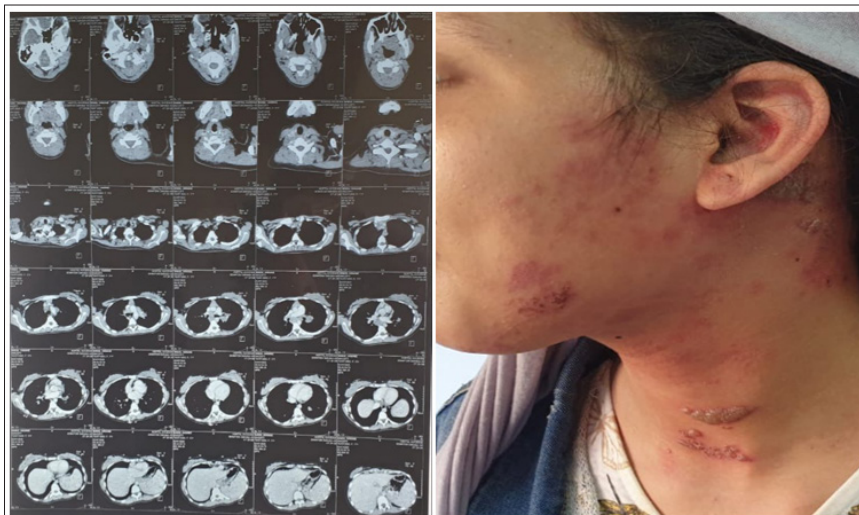


Figure 1: Cervical-thoracoabdominal axial CT scan

Figure 2: Clinical photograph showing vesicular lesions

Discussion

Infection of the human body with VZV consists of three phases starting with the primary infection where the virus enters the body through the lining of the upper airways (nasopharynx) and oropharynx. It multiplies and then spreads in the body by a first viremia. The virus replicates locally in the respiratory tract and the lymph nodes, then reaches the reticuloendothelial system. The specific immune response develops, a second viremia disseminates the virus in the whole organism and brings it to the level of the skin (vesicular lesion appears), the latency phase follows the recovery of varicella. It is characterized by the persistence of viruses in the sensory ganglia of certain spinal nerves and or cranial and settles in a quiescent state.

The mechanism by which the virus remains in the latent state is very imperfectly known [5]. In this phase, there is no clinical manifestation. The virus reactivation phase occurring due to immune deficiency is common in elderly subjects and in pathological situations affecting the immune system [6,7]. Such as in our case, tuberculosis, which leads to a decrease of immunity. It is transmitted by the sensory nerve to the epidermis to give localized shingles infection.

Concerning the relation between herpes zoster and tuberculosis, ARVID WALLGREN concluded that tuberculosis in one way or another predisposes to herpes zoster by the action of its toxins on the spinal ganglia. In a group of 10 patients, 7 cases present with tuberculosis disease or infection, and the remaining 3 cases which were tested for tuberculin were not attempted [8]. The frequency of herpes zoster in tuberculosis patients was demonstrated in 21 autopsies. HEAD and CAMLBELL encountered 5 times extensive pulmonary tuberculosis and 5 times milder tuberculosis lesions [9].

ARNSTEIN finds that in half of the cases where the autopsy reveals an alteration of internal organs, there is a combination of herpes zoster with tuberculosis. He believes that the chronic irritation of the spinal ganglia by tuberculous toxins promotes the outbreak of disease [10]. MAGNUSSON, from REYKAVIK further expresses the opinion that tuberculosis or tuberculosis

toxins can give rise to herpes zoster infection [11]. Furthermore, ARVID WALGREN believes that the herpes zoster pattern corresponding to the location of the tuberculous lesions shows that the majority of his patients had the same site of herpes zoster as the tuberculous lesions.

Conclusion

The association of herpes zoster infection and tuberculosis has already been mentioned in some studies; moreover, its cervicofacial location and a lymph node form of tuberculosis have been reported in our case. Many studies have described Herpes zoster infection complicating tuberculosis. However, and to the best of our knowledge, an association of cervicofacial herpes zoster and lymph node form of tuberculosis hasn't been reported yet in the literature, hence the rarity of our case.

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