

Case Report

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Idiopathic Thrombocytopenic Purpura in Tuberculosis

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ABSTRACT

Tuberculosis (TB) is a contagious disease in both developing and developed countries. The incidence is increasing due to bacilli resistant to many drugs and the human immunodeficiency virus (HIV). Since 2012, India has the highest incidence of the disease at around 2.2 million cases, accounting for 26% of the global incidence according to World Health Organization statistics. A wide range of hematological manifestations is observed in TB, where thrombocytopenia is common in miliary TB and thrombocytosis in pulmonary TB. Immune thrombocytopenic purpura (ITP) is characterized by a low platelet count accompanied by the presence of autoantibodies. The association of immune thrombocytopenic purpura and tuberculosis is a rare condition. In the case presented, anti-tuberculosis therapy was effective for both tuberculosis and thrombocytopenia, suggesting a causal link between tuberculosis and immune thrombocytopenic purpura.

Purpose: Reporting a case of association of PTI and TBC

Method: Clinical case study and correlation with literature

Discussion: The pathophysiology of thrombocytopenia in tuberculosis remains unanswered. This is a rare condition, estimated to occur in less than 1% of tuberculosis cases. Mycobacterium tuberculosis can break down the antigen with platelets leading to the formation of antiplatelet antibodies. HLA-specific presentation of tuberculosis may also lead to antiplatelet immune response in some patients.

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Patient S.SH, male, age 64 years, born Korca (known endemic area for TBC) is referred by the district hematologist as suspect immune thrombocytopenia.

The patient's complaints were related to fatigue, weakness, sweating, blood in the urine, blood through saliva and black stools, which the patient had been complaining about for almost 4-5 days. Objective examination: Pale on the skin and mucous membranes with obvious signs of cutaneous hemorrhage in the form of diffuse ecchymoses and petechiae, dense, widespread throughout the body but more evident in the upper and lower bilateral extremities as well as at the level of the right hemithorax and abdomen. Patient with signs of urinary tract hemorrhage (evident in the catheter bag) as well as bleeding gums all the time.

Clear conscience, rhythmic cardiac tone with frequency 85bpm, TA = 100 / 80mmHg, attenuated respiration more expressed on the right side, Sat O₂ 96% under room air, soft abdomen. Treatable not dolent on palpation abdomen, without muscle protection, liver and spleen below the costal arch, no peripheral palpable lymphadenopathy.

Anamnesis Vitae

Patient Known as Diabetes Mellitus type 2 under treatment with

oral antidiabetics, HTA under treatment with antihypertensives.

Family History: Nothing to note

Examinations Shown

Hemograms RBC 3.41x10⁶, Hb 9.0gr / dl, Hct 26.9%, MCV 79.0fL, RDW 14.6%, WBC 8.2K / uL, PLT 2K / uL (norm 150-400). Thrombocytes in the slide: rare in preparations. Reticulocyte count 23.0% / °°

Bone Marrow Aspiration

Promyelocytes 1%, Neutrophil myelocytes 2%, Neutrophil metamyelocytes 8%, Neutrophil sticks 22%, Segments 25%, Lymphocytes 13%, Monocytes 7%, Polychromatophilic normoblasts 12%, Oxyphilic normoblasts 8%, Plasma cell 2%. rich in material, M / E ratio = 2.9 / 1. Megakaryocyte apparatus present with a small platelet burst, Leukocyte immunophenotype: Bone marrow material. Through leukocyte immunophenotyping was observed:

About 1% of cells with the markers CD34 +, CD117 +, HLA-DR +, CD13 +, CD33 + probably myeloblasts

1. Lymphoid population makes up about 6% of all cells which were studied for the above markers
2. Granular series which makes up about 79% of all cells parasites with normal granularity
3. Monocyte population makes up about 4% of all cells
4. The red series makes up about 11% of all cells

Coagulation Panel

PT 72%, INR 1.24, aPTT 23.0sec, Fibrinogen 471mg / dL, D-dimer 0.82µg / mL

Biochemical Balance

Glucose 157mg / dL, Urea 124.2mg / dL, Creatinine 1.18mg / dL, Direct bilirubin 0.33mg / dL, ALT 11UI / L, AST 16U / L, GGT 28U / L, APL 50U / L, LDH 259U / L, CK 132U / L, Calcium 8.8mg / dL, CRP 1.46mg / dL.

Immune Panel

HBsAg0.30S / CO (<0.99 neg) Anti-HCV 0.88S / CO (<0.99 neg), Anti-HBs 0.09IU / L (<9.99 neg), AntiHBc 0.17S / CO (<1 neg), HIV 1 & 2 Ab, RDT negative, ANA <160 (negative), Anti ds DNA 8.4IU / mL (<35, neg) ENA 0.10U / mL (<0.8 neg), MPO 2.9U / mL (<10 neg).

CEA 0.8 (<10 neg), AFP 3.8 (<8.8 neg), CA 19-9 10.4 (<37 neg), CA 15-3 9.9U / mL (<31.3 neg), Total PSA 0.43ng / mL, Ferritine 130.70 ng / mL.

Lung CT

Trachea, free main bronchi. Sinister pleural thickening with posterobasal effusion with DIP 16mm. Multiple pseudonodular consolidations in both lungs.

Protein Electrophoresis

Albumin 58.5%, alpha 1-globulinate 5.3%, alpha 2-globulinate 13%, beta 1-globulinate 5.5%, beta 2-globulinate 4.3%, gammaglobulinate 13.4%, Ratio A / G 1.41.

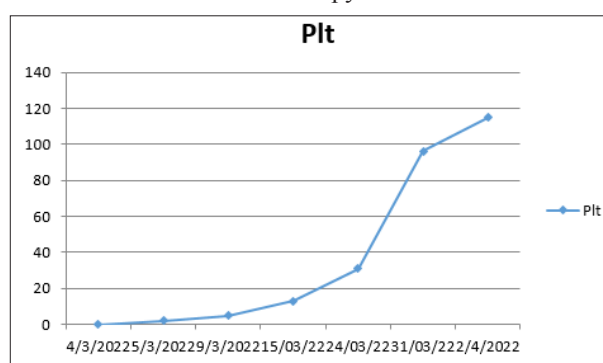
After completion with the necessary examination panels the patient was placed under corticosteroid therapy with a maximum dose of 2 mg / kg body weight.

On 08/03/2022 the patient is consulted by a pulmonologist who requires equipment with Sput-BK 3 samples which Geneexpert sputum (Positive) showed the presence of DNA for MTB positive.

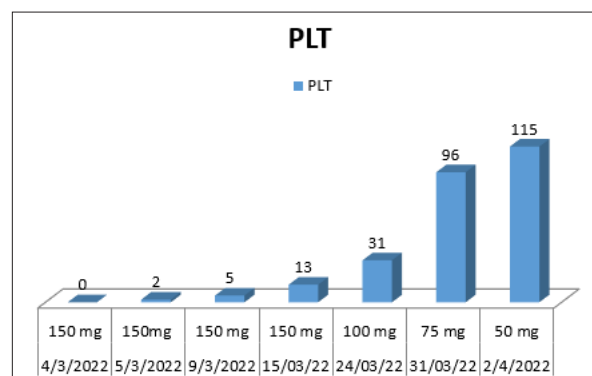
The Patient Was Placed from 11/3/2022 under the Anti-Tuberculosis Regime with

1. Rifampicine / Isoniazid 150/75 mg 3tb per day oral
2. Pyrazinamide 500 mg 2tb per day oral
3. Gambutol 400 mg 2 tb per day oral
4. Multivitamine 3x1 tb per day oral

The Following is the Course of Platelet Counts Before and after the Start of Antituberculous Therapy



The Graph below Shows the Correlation between the Doses of Cortisone used by the Patient and the Value of PLT. The Patient from 11/03/2022 is under the Regime with three Anti-Tuberculosis



Discussion

TB is ranked as the 13th leading cause of death in the world and the second leading killer of infectious origin after the COVID-19 period (on HIV / AIDS). In 2020 about 10 million people were infected with tuberculosis according to WHO reports of which 5.6 million men, 3.3 million women and 1.1 million children. TB is present in all countries and age groups.

The disease was under control especially in developing countries, until the emergence of HIV infection and the identification of resistant strains of mycobacteria [1-3]. Various hematological manifestations include abnormalities such as anemia, leukocytosis, monocytosis, lymphopenia, leukopenia, thrombocytopenia, thrombocytosis, leukomoid reactions or even pancytopenia , but the thrombocytopenia is deep in tuberculopathy and tuberculopathy there are some reports (in total 18 cases reported) [4-10].

Idiopathic thrombocytopenic purpura (ITP, also known as primary immune thrombocytopenic purpura) is an acquired disease that is defined as isolated thrombocytopenia without other potential causes present. It usually has a secret onset with a long history of purpura (purpura). > 6 months), spontaneous remission is uncommon and in most cases is likely to be incomplete [11]. Steroids are conventional first-line therapy for adult ITP. Platelet count increases in responding patients and usually reaches maximum values from two to four weeks after the start of treatment. However, in most patients, the platelet count drops when the steroid dose is reduced or discontinued. Another alternative treatment for primary autoimmune TPI is the use of venous immunoglobulins, the use of which is indicated in cases where certain clinical situations require an increase in immediate due to interventions or clinically significant complications [11]. In our case, the exclusion of ITP was done not only based on standard criteria, but also with the fact of the reaction of platelet value after the combination of antituberculous therapy accompanied by antituberculous therapy. Other causes of thrombocytopenia such as hemophagocytic syndrome, TTP, combined autoimmune clinical and laboratory cytopenia, bone marrow aspiration, and biopsy described in the case report were also excluded.

It is already known that various factors cause hemorrhage related to infectious conditions, of which the most common is thrombocytopenia. The etiology of thrombocytopenia in most cases appears to be an increase in platelet destruction, either due to KID or septicemia, as well as platelet adhesion to damaged vascular surfaces, or direct platelet toxicity caused by other microorganisms from the involvement of the bone marrow in the infectious process. Although the most important therapy for Infection-Related Thrombocytopenia is that directed at the underlying infection, treatment decisions for thrombocytopenic purpura immunity remain controversial and may include single or combination

therapy with corticosteroids, intravenous immunoglobulin (IV). thrombocytopenia or hemorrhage [11]. In the reported case the clinic was dominated by gingivorrhage, hemoptysis, diffuse petechiae and ecchymoses, macroscopic hematuria. Based on clinical examinations, radiological findings (chest x-ray and CT), demonstration of acid-fast bacilli positivity in sputum (Gene expert) and after excluding other causes of thrombocytopenia, the diagnosis of purulent immune thrombocytopenia due to TBC was made tuberculosis.

The patient was placed on corticosteroid therapy and from the follow-up to the dynamics no significant response to platelet count was observed, until the moments after the start of antituberculous therapy when the improvement of values was proven not only in laboratory but also with improved clinical performance [11].

The effect of steroids on thrombocytopenia is complex and has a late effect. The mechanism of action of intravenous immunoglobulins is unclear, but studies suggest blockade of Fc receptors in the cellular reticuloendothelial system and inhibition of antibody production and binding, thought to be associated with anti-idiotypic antibodies that bind to antithrombotic antibodies.

In our case, the corticosteroids were discontinued on the 14th day of therapy and the patient was discharged from the hospital and thrombocytopenia was not detected after discontinuation of corticosteroid therapy.

These observations suggest that tuberculosis is the cause of thrombocytopenia in our patient. He had no hemolysis present, hemophagocytic syndromes, gastrointestinal hemorrhage or in other regions. Thirteen days after discharge from the hospital, the patient was clinically healthy with a platelet count of $300 \times 10^9 / l$. The patient is still in our follow-up without clinical or laboratory findings confirming the recurrence of thrombocytopenia.

In cases where TB cases reported by the WHO are present in every corner of the world and since TB is defined as the second most common infectious cause of death in the world, the thrombocytopenic purpura in TB should remain the focus of clinicians. To clarify the exact immune mechanisms more detailed studies are needed, and they will probably enable even innovative therapies related to immune thrombocytopenia in TBC making it possible to eliminate possible side effects of corticosteroids.

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