

BILATERAL UPPER AND LOWER LIMBS MUSCULOSKELETAL AMYLOIDOSIS WITH OSTENSIBLE POLYARTHRITIS IN YOUNG ADULT – A CASE REPORT

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Abstract

Background: Rheumatoid Arthritis is a chronic, autoimmune disorder marked by damaging synovial tissues. It is associated with erosive alterations and particularly affects tiny joints in the body. **Case presentation:** A 19-year-old female adult with ostensible polyarthritis was admitted to the outpatient department. She had bilateral upper and lower limbs musculoskeletal amyloidosis. Her symptoms worsened during the COVID period and continued for one year. Her initial treatment started with glucocorticoids (GC) and methotrexate (MTX). But her symptoms developed. Finally, the treatment with Tab Dexamethasone (4mg), Tab Zerodol, Tab Folvite (5mg), and Tab HCOs (200mg) continued, and the patient was put on a regular follow-up schedule. The ESR value decreased (52mm/hr) and the patient's condition improved significantly. **Discussion:** Younger adults with rheumatoid arthritis are dealt with different issues either the disease or the treatment modalities. In this case, a variety of steroid-sparing agents have been employed. **Conclusion:** To manage rheumatoid arthritis in young adults, special consideration is required. Providing adequate education to the patient and appropriate follow-ups could prevent further functional disabilities of musculoskeletal disease.

Keywords: Rheumatoid Arthritis, Musculoskeletal Amyloidosis, Bilateral Limbs, Young Adult, Ostensible Polyarthritis.

INTRODUCTION

Rheumatoid Arthritis (RA) is a systematic autoimmune and inflammatory disorder primarily affecting synovial tissue. It leads to joint destruction and disability. In 2019, 18 million people worldwide were living with rheumatoid arthritis [1]. Approximately 1% of the world's population suffers from RA. Joint pain and swelling with a limited range of motion is a common manifestation of RA. It is a heterogeneous disease that encompasses different subtypes of arthritis, depending on the presence of the enteritis and the number of affected joints [2]. RA patients have an extra-articular manifestation of rheumatoid vasculitis (RV), which started to evolve in the 1960s. The clinical manifestations of vasculitis include skin rashes, cutaneous ulcerations, and visceral infarctions. Musculoskeletal amyloidosis (MA) is a complex disease process. It tends to be systemic but frequently poses a diagnostic challenge [3]. Amyloidosis is classified into many types. The four types of systemic amyloidosis are primary arthritis, familial arthritis, wild-type systemic arthritis, and secondary amyloidosis. In RA,

secondary arthritis was the most common type of systemic arthritis. Amyloid deposition presents with a variety of symptoms such as liver involvement, restrictive cardiomyopathy, and nephrotic syndrome [4]. It also presents in peripheral nerves and the gastrointestinal tract. This study is of a 19-year-old young adult with bilateral upper and lower limb musculoskeletal amyloid with ostensible polyarthritis initially diagnosed and managed as a case of RA.

CASE PRESENTATION

A 19-year-old female adult with ostensible polyarthritis was admitted to the outpatient department in a tertiary hospital in Chennai City. She had pain in several small joints in her body for five years. She also presented with generalized bone pain, swelling, and contractures of both knees. On examination, she had bilateral upper and lower limbs musculoskeletal amyloidosis. Her symptoms worsened during the COVID period and continued for one year and were not associated with trauma. Soon she developed a severe morning stiffness lasting up to 2 hours. She was born into a family of non-consanguine parents. Her family history revealed that her relative was affected by RA and was treated for 5 years with methotrexate and rituximab. The laboratory result showed persistently elevated inflammatory markers with a negative antinuclear antibody (ANA) screen. Her initial treatment started with glucocorticoids (GC) and methotrexate (MTX). But her symptoms developed and affected wrists, elbows, ankles, small joints of both hands and legs, and severe uveitis of the right eye. Then a good response was felt after using biologic therapy with infliximab. After a few months, adalimumab was used due to relapses. Again, some joints had a remission characterized by swelling and pain. ESR (Erythrocyte Sedimentation Rate) and CRP (C - reactive protein) were measured. ESR was critically high (119mm/hr) and CRP was negative (7.7mg/dl). During the medication therapy, Tab Zerodol and Tab Wysolone(10 mg) were used and she depended on a daily dose of Tab Rantac and Inj Dexamethasone (100mg). Gradually she had a low bone mineral density, regardless of the vitamin D.



Figure 1 [A] MRI scan showing multiple cross-sectional views [B] MRI scan shows a horizontal slice, with the front (anterior) part of the body at the top of each image and the back (posterior) at the bottom. [C] MRI scan of the knee, which provides a more detailed view of the soft tissues, cartilage, and other structures within the knee joint compared to X-rays

Her investigations showed hemoglobin of 8 g/dl. She received cyclophosphamide (6x) and rituximab (3x). Patient's elbows, shoulders, radiocarpal joints, metacarpophalangeal, distal interphalangeal joints, and both knees had a restricted range of movements. The treatment with Janus kinase (JAK) inhibitor tofacitinib was initiated, which led to the gradual amelioration of musculoskeletal symptoms. She had many consequences of the disease by prolonged GC treatment. Finally, the treatment with Tab Dexamethasone (4mg), Tab Zerodol, Tab Folvite(5mg) and Tab HCOs(200mg) continued, and the patient was put on a regular follow-up schedule. The ESR value decreased (52mm/hr) and the patient's condition had improved significantly.

DISCUSSION



Figure 2: [a] X-ray of the anterior-posterior (AP) view of both knees in a standing position [b] bilateral knee X-ray taken from the lateral (side) view. The X-ray shows the right and left knees of a patient showing the femur (thigh bone), tibia (shin bone), and patella (kneecap).

RA is a chronic, autoimmune disorder marked by damaging synovial tissues. It is associated with erosive alterations and particularly affects tiny joints in the body. It also causes low quality of life and diminished functional ability of the patients [5]. Younger adults with rheumatoid arthritis are dealt with different issues either the disease or the treatment modalities. In this case, a variety of steroid-sparing agents have been employed [6]. The use of the tofacitinib study in patients showed improvement in symptoms and functional disability. Tofacitinib is an oral agent that younger adults should not overlook. Different diagnosis was considered in younger adult patients along with various treatment modalities. This data will be beneficial for physicians as they consider the positives and negatives of treatments. X-rays of the bilateral knee and feet are useful to view changes in inflammatory arthritis [7]. Several clinical trials of rheumatoid arthritis in adult patients have given evidence about the use of tofacitinib. According to 3 case reports of rheumatoid arthritis [8,9,10] cardiac involvement is the leading cause of organ injury affecting patients with amyloidosis.

CONCLUSION

This study portrays a rare occurrence of bilateral upper and lower limbs RA in younger adults that is rarely reported in the medical literature. To manage rheumatoid arthritis in young adults, special consideration is required [11,12]. Providing adequate

education to the patient and appropriate follow-ups could prevent further functional disabilities of musculoskeletal disease [13]. This case report emphasizes the challenges in the management of young adults with long-term musculoskeletal disease and treatment modalities.

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Conflict of interest

Authors of this article have no conflict of interest.

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