Staphylococcus Aureus Thigh Abscess in A Systemic Lupus Erythomatosus Young Adolescent Patient

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Abstract: Systemic lupus erythematosus (SLE) rarely present as a soft tissue localized abscess. Occurrence is further rare in young adolescent age as there are finger countable cases in the world literature. The abscess was treated with combination of antibiotics and surgical drainage successfully.

Keywords:
1. Case Report

A 19 yr Old girl was hospitalized with the diagnosis of SLE based on presence of Butterfly rash, mouth ulcers, photosensitivity, proteinuria, Antinuclear antibodies (nuclear homogenous pattern), dsDNA antibodies, arthralgias of small and large joints and anemia (Hb 9.2 gm%) fulfilling the ACR classification criteria for SLE, 7/11. There was no previous history of medication use, radiation, drug abuse, surgery trauma or infection. She was under treatment as a case of SLE with prednisolone 2mg/kg/day. Initially, she was on Hydroxychloroquine, the dose of steroids was tapered to 15 mg alternate day which the patient was receiving on her hospitalization. She did not receive Azathioprine/ cyclophosphamide. She offered history of insidious pain in the middle of right thigh without cramps and fever or apparent trauma past 4 days. The physical examination revealed mouth ulcers, malar rashes, stomatitis, joint pains, localized muscle tenderness and woody indurations. However, there was no rise in local temperature or oedema. The laboratory test revealed Hb 9.2 gm%, HCT 27.1%, WBC 6400 / cumm (85% neutrophils, 12% lymphocytes and 2% monocytes), Platelet count 1, 71000/ cmm. ESR 52 mm first hour, proteinuria ++, urea 25 mg/dl and creatinine 0.9 mg/dl. The muscle enzymes tests were normal. Alanine aspartate, creatinine kinase and amino transferase were within normal limits. ANA and anti dsDNA was positive. The high frequency ultrasonography of right side thigh lateral medial aspect showed collection of 350 cc in between vastus medialis and gracialis suggestive of an abscess( Figure-1). Blood culture did not identified any organisms. As the pain in the middle thigh was excruciating patient was subjected to incision and drainage of abscess and 300 cc could be drained. The pus culture revealed staphylococcus aurius. Treatment with IV cefuroxime 1.5 gm BD and metronidazole 500 mg 8 hry was administered for 7 days. She was discharged after one week on oral steroids in tapering doses of prednisolone 0.5 mg/kg/day with cyclophosphamide 2mg/ kg/ day with lincomycin for 7 days. Patient had a sound recovery after clinical treatment.
Legend to figure (1)- High frequency ultrasonography of Lateral medial aspect of right thigh showing elongated collection (pus) seen in between vastus medialis and Gracilis.

2. Discussion

SLE is one of the diseases thought to predispose to several severe infections like salmonella, Nocardia, tuberculosis, staphylococcus aureus, streptococci species & E.coli[1, 2, 3, 4]. SLE patients have an increased risk of infections due to immunological dysfunction and use of chemotherapy and immunosuppressive agents. Despite increase propensity of SLE patients to develop opportunistic infections, the muscle infection – pyomyositis and localized muscle abscess are rarely observed with the disorder. The presence of localized pyomyositis or abscesses appear to be considered a vital feature during the initial differential diagnosis. The delay in diagnosis may result in increased significant morbidity and sometimes mortality. The present patient had thigh abscess caused by staphylococcus aureus. The antibiotics were given inaccordance to sensitivity report. Tung Chen Y and Isenberg D described necrotizing fasciitis in SLE and describe 32 cases, staphylococcus aureus was noted in only 1 case.[5] Blay G et al in 2014 reported a case of purulent stage of pyomyositis in the middle third of vastus medialis without elevation of muscle enzyme where in computer tomography of left thigh showed collection of pus. The author claims the case as first ever described in the lupus population in pediatric age group in april 2014. The present case is similar to one described by Blay G et al who presented as a case of SLE as per American college of rheumatology ACR classification criteria[6]. The patient presented with localized severe muscle pain with induration in the right middle aspect of thigh. On ultrasonography the lesion revealed elongated collection in intramuscular plane in medial aspect of right upper thigh between vastus medialis and Gracilis. The drainage with the aid of ultrasound yielded 300 cc frank pus. On microscopy contained abundant pus cells and leukocytes. The pus culture revealed staphylococcus aurius. The muscle enzymes were not elevated in the case.

The infectious myopathy was first described in 1885 by Scribbas[7], with most affected regions are thighs and hips. Review of data from jan 1983 to july 2013 revealed only one case (0.34%) with pyomyositis episode with SLE[8]. Similar clinical features may also occur in healthy patients after local trauma and may be associated as non specific pyomyositis with associated diseases like leukemia, diabetes, scleroderma, HIV, renal disorders. However, to the best of our knowledge and review of world literature, occurrence of pyomyositis / localized abscess in young adolescent lupus cases remains a rarity. The differential diagnosis of such localized pyomyositis affecting thigh includes deep venous thrombosis, hematoma, cellulitis, osteomyelitis etc. Intravenous broad spectrum antibiotics and per cutaneous surgical incision and drainage remains the main orchestra of treatment besides primary management of the underlying disease. Our patient had complete recovery after clinical treatment.

3. Conclusion

A case of staphylococcus abscess in right middle thigh of a young adolescent patient with SLE is described. The case emphasizes the importance of paying due attention to localized muscle pain in immune compromised patient even without the elevation of muscle enzymes which raises the suspicion of underlying abscess and demands ultra ultrasonographic or CT imaging in the case. A prompt antibiotic therapy is warranted besides incision and drainage. The muscle infection is rarity in this adolescent young age group who are presenting with SLE.

References