

# A Fatal Presentation of Dermatomyositis

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## ABSTRACT

Dermatomyositis (DM) is an autoimmune disease that mainly affects the skin, muscle, and lung. The pathogenesis of skin inflammation in DM is not well understood. We present a 40-year-old male who presented with four months history of generalized skin lesion, pain and weakness. He had elevated transaminases with detectable muscle weakness. This case highlights the need to consider dermatomyositis with skin lesion and weakness; and the use of aggressive immunosuppressive therapies due to its associated vasculopathies.

## KEY WORDS

*Cutaneous lesions, dermatomyositis, idiopathic, myopathy, vasculitis*

## INTRODUCTION

Dermatomyositis (DM) is a chronic inflammatory disorder that can affect the skin, muscle, and other organs and is associated with significant morbidity and mortality.<sup>1-6</sup> The prevalence of DM is not well-defined, as it is historically grouped together with polymyositis (PM) and inclusion body myositis (IBM) in most epidemiologic studies.<sup>7</sup> Currently classified as an idiopathic inflammatory myopathy (IIM), much of the work in understanding DM has been focused on the muscle pathology that accompanies this disorder.<sup>8</sup> The inflammatory mechanism is attributed to a primary T-helper-cell dependent, B-cell-mediated, local humoral immune mechanism, which causes vascular occlusion and capillary obliteration with ischemic change in the skin and muscle.<sup>4,9,10</sup> Dermatomyositis may be associated with systemic manifestations including restrictive and interstitial lung disease and cardiomyopathy.<sup>4</sup> We present a 40-year-old male who presented with four months history of generalized skin lesion, pain and weakness.

## CASE REPORT

A 40 years male from Devitar presented to Kathmandu University Hospital (Dhulikhel Hospital) in July 2011 for evaluation of generalized itching for six months; pain, weakness and skin lesions in different part of body for four months. The onset of the disease was acute, gradually progressive; he could eat with his hands but couldn't brush his teeth, could wear slippers but couldn't walk upstairs. This weakness amplified during the 15 days before his presentation. For the same duration gives history of scalp hair loss. He had some antibiotic medication for this, the nature of which could not be assessed. Although there was a history of subjective fever, there was no history of oral ulcers, shortness of breath, or recent immunizations. He was a farmer by occupation, led an active working lifestyle, non-smoker and occasional alcoholic. His medical and family history gave no abnormal findings.

At the time of admission, on examination, general condition

**Table 1. Laboratory values - at presentation, and discharge (after three weeks).**

	Normal Range	At presentation	At discharge
Sodium	134–143 mmol/L	135	130
Potassium	3.5–5.0 mmol/L	4.2	4.1
Lactate dehydrogenase	470–750 units/L	6310	-
Aspartate aminotransferase (SGOT)	15–40 units/L	137	-
Alanine aminotransferase (SGPT)	10–55 units/L	174	-
Alkaline phosphatase	178–455 units/L	127	-
Urea Nitrogen	7–17 mg/dL	36	-
Creatinine	0.20–0.70 mg/dL	0.6	-
Bilirubin total	0.2–1.2 mg/dL	1.2	-
Bilirubin direct	0.1–0.2 mg/dL	0.15	-
Serum glucose	60–115 mg/dL	63	74
Creatine phosphokinase (CPK)	30–150 units/L	1879u/L after 4 days of admission 9980u/L after 9 days of admission 17500u/L after 16 days of admission	
Erythrocyte sedimentation rate	0–13 mm/h	22	-
WBC counts (N-85%,L-11%,E-3%)	5.0–13.0 x 10 <sup>3</sup> /mL	12.3	10.4

was ill looking, height - 156.5 cm, body weight 57.5 kg, and body temperature 37.2°C. blood pressure was 119/ 78 mmHg, pulse rate 73 per minute with a regular rhythm, and respiration 20 breaths per minute; liver palpable 5 cm below right sub-costal margin (RSCL). Cardio-pulmonary examinations were normal, neurological examination shows power on proximal muscle as 0/5 whereas distal muscle showed 5/5 both on upper and lower limbs. Sensation was intact, with normal bowel and bladder habit.

Skin changes include pigmentation over malar space, reticulate pigmentation over abdomen, poikiloderma over lower limbs, Gottron's sign, Shawl sign, Holster sign, vasculitis present over distal fingers, tenderness over digits. Typical erythematous lesion was not noticed in this patient, which could be because of the darker skin color of the patient.

Laboratory investigations are shown in table 1. The abnormalities include increased CPK of 17500u/l on 16th day of admission, raised Lactate dehydrogenase of 6310 units/L, elevated WBC count of 12.3 x 10<sup>3</sup>/mL (N-85%,L-11%,E-3%), elevated Aspartate aminotransferase of 137 units/L and elevated Alanine aminotransferase (SGPT) of 174 Units/L. Ultrasonography presented fatty hepatomegaly. Muscle Biopsy indicated mild to moderate chronic inflammatory cells, presence of perivascular inflammatory cell, few atrophic and degenerative fibers suggesting active necrotizing myopathy, consistent with dermatomyositis.

However ANA and dsDNA were found to be negative. On third day of admission patient developed dysphagia but denied any nasogastric feed. He had no other neurological impairment. On the tenth day of admission he developed shortness of breath and fever with chest examination revealing crepitation over bilateral lung field.

He was treated with prednisolone 60mg initially then 80mg. Antibiotics- Amoxiclav and metronidazole were used for respiratory tract infection. He also underwent physiotherapy and psychiatric consultation was done as the patient seemed as depressive.

By third week, patient was improving on his muscle weakness; with muscle power 3/5 on proximal upper and lower limbs. The respiratory tract was also improving with decrease in crepitation. The patient was discharged on demand for Hindu festival of Dashain/Dashara. The patient was asked for follow but we were sad to hear that the patient expired after five days of being discharged from the hospital with the possible expected cause of death being aspiration pneumonia. Post mortem was not done, during this festival holiday.

## DISCUSSION

Dermatomyositis (DM) is a systemic inflammatory disorder affecting the skeletal muscles, the skin, and other organs.<sup>11-13</sup> The diagnosis of DM is still based on Bohan and Peter criteria: 1) symmetric proximal muscle weakness; 2) increased serum muscle enzymes; 3) myopathic changes upon electromyography; 4) typical histological findings on muscle biopsy; and 5) typical dermatologic manifestations, such as heliotrope rash or Gottron's papules.<sup>4,14</sup> Diagnosed positive if any 4 out of 5 of the above criteria met, although latter criteria are most often used, they have several limitations (e.g., they do not include myositis-specific autoantibodies). We could meet four of the criteria for diagnosing presented case.

The estimated incidence of DM is approximately 1 per 100,000 per year.<sup>7</sup> There are two forms of DM, juvenile and adult, those have overlapping but some distinct clinical features.<sup>7</sup> There is a female to male predominance of about



Figure 1. Gottron's sign in dermatomyositis.



Figure 2. Gottron's sign(legs) in dermatomyositis.



Figure 3. Facial erythema (erythroderma) in a patients with dermatomyositis.



Figure 4. Heliotrope rash in dermatomyositis.



Figure 5. The shawl sign in dermatomyositis.

2:1. The peak incidence in adults occurs between the ages of 40 and 50, but individuals of any age may be affected.<sup>15</sup>

Histologic features of DM include muscle fiber necrosis, degeneration and regeneration, and an inflammatory cell infiltrate. DM is considered to be a humorally mediated disorder in which the cellular infiltrate, located principally in perifascicular regions, is often focused around blood vessels.<sup>17-19</sup> The terminal complement C5b-9 membrane attack complex is detectable in vessel walls before the appearance of inflammatory cell infiltration in DM. This is absent in Polymyositis. The inflammatory infiltrate is composed of B cells and plasmacytoid dendritic cells that are CD4+. Other typical features include perifascicular atrophy and fibrosis. Abnormal muscle fibers are usually grouped in one portion of the fascicle, suggestive of microinfarction mediated by blood vessel dysfunction. (refer comment 12). The early activation of the complement cascade leads to the formation and deposition of the membranolytic attack complex on endomysial capillaries,

resulting in their destruction and tissue ischemia.<sup>17,20</sup>

Although this key pathophysiologic event has strengthened the prospects for targeted immunotherapy in these conditions, the lack of identification of target auto antigens represents a major hurdle for the definition of specific immunotherapy in Dermatomyositis. Nevertheless, to date, nonspecific immunotherapeutic drugs have markedly improved the outcome of DM patients, and treatment does not selectively target pathogenic auto reactive T cells in PM or complement-mediated processes in DM.

Dermatomyositis is a multisystem disorder with a wide variety of potential clinical findings. Muscle weakness is the most common presenting feature of DM. The onset is usually insidious, with gradual worsening over a period of several months before medical attention is sought, very similar to this case presentation. The distribution of weakness is typically symmetric and proximal. An inability to swallow and symptoms of aspiration can reflect involvement of striated muscle in the pharynx or upper

esophagus; dysphagia or dysphonia is generally associated with a rapidly progressive course.<sup>20,21</sup> Several distinct rashes, generally present at the time of clinical presentation, occur in Dermatomyositis- Gottron's sign, Heliotrope rash, Shawl sign and V sign, Erythroderma, Periungual abnormalities, Psoriasiform changes in scalp. DM may affect the lungs primarily, resulting in interstitial lung disease (ILD), which may lead to life threatening complications (i.e., ventilatory failure, secondary pulmonary arterial hypertension, or cor pulmonale).<sup>22</sup>

Esophageal motor activity disorders related to PM/DM may be responsible for swallowing disorders, including difficulty with solids and liquids, pain and pre-prandial discomfort in the sternal area, gastro esophageal reflux into the pharynx and/or mouth, coughing while eating, as well as life-threatening complications (e.g., aphagia to solids/liquids requiring total enteral feeding and aspiration pneumonia).<sup>12,21-24</sup> Cardiac impairment may occur in up to 50% of patients, although it is more often asymptomatic in DM.<sup>1,25</sup> Various abnormalities have been described, including mainly conduction defects and primarily end-rhythm disturbances; congestive heart failure, pericarditis, and valvular disease are severe, less frequent cardiac complications in PM/DM patients.<sup>1,25</sup> Evidence for an association between PM/DM and solid tumors came first from case reports, with the reported rate of cancer varying from 6% to 60%.<sup>1,12,23,26-30</sup>

Lab investigation shows raised muscle enzyme - creatine kinase, lactate dehydrogenase, aldolase, and aminotransferases; and serological test can be ANA positive, Anti Jo positive. Autoantibodies are found in a majority of patients. Antinuclear antibodies in high titer suggest the presence of another connective tissue disease. Skin and/or muscle biopsies may establish the diagnosis in DM. Biopsy of a variety of DM skin findings, including Gottron's sign, the shawl sign, and erythroderma can provide confirmation of the diagnosis in the proper clinical setting. Histology of muscle shows perifascicular, perimysial, or perivascular infiltrates, perifascicular atrophy. MRI, chest X-ray, Barium

swallow can also be done, to see involvement of other organ system and MRI shows extent of damage in muscle.

Treatment for myopathy is done with high dose of corticosteroid. In our case, we started with 60 mg of corticosteroid and later dose was increased to 80 mg. Alternative drugs that can be used are *Azathioprine*, *Methotrexate*, *Mycophenolate mofetil*, *Monoclonal anti-CD20 (Rituximab)*, *Cyclophosphamide*, *Immunomodulation with Ivlg*. Our strategy was slow change of treatment to *Azathiorine*, on follow up schedules. Treatment for skin lesion requires avoidance of sun exposure with immunosuppressant like *Methotrexate*, *Mycophenolate mofetil*, *Hydroxychloroquine*, and *Chloroquine*. Extra protein diet and observing the patient for development of complication is a factor to be considered in treatment.

## CONCLUSION

Dermatomyositis is a rare disease entity. Prognosis depends on severity of myositis, presence of malignancy, cardiopulmonary system involvement. Patients severely affected at presentation or treated after long delays, those with severe dysphagia or respiratory difficulties, older patients, and those with associated cancer have a worse prognosis. The treatment of Dermatomyositis includes high-dose glucocorticoids in addition to disease-modifying agents or cytotoxic agents such as methotrexate. Most patients improve with therapy, and many make a full functional recovery, which is often sustained with maintenance therapy. Up to 30% may be left with some residual muscle weakness. Relapses may occur at any time. Early management of disease, diagnosis of any internal malignancy, any toxicity of treatment is important to look on follow up.

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