A case of complicated bullous systemic lupus erythematosus managed with Rituximab
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ABSTRACT

Skin involvement is seen in nearly 76 per cent of Systemic lupus erythematosus (SLE) patients, but bullous lesion however accounts for less than one per cent of cutaneous manifestation of lupus. Bullous systemic lupus erythematosus (BSLE) is a distinctive entity characterised by histological appearance of sub epidermal bulla with deposition of pan immunoreactive determinants and circulating antibody against type VII collagen (NC1 domain) and occasionally other antigens like laminin 5, laminin 6, and bullous pemphigoid antigen (BP230). BSLE frequently parallels internal organ involvement. In this case report we have reported a case of bullous systemic lupus erythematosus (BSLE) who presented with an organ threatened condition in the form of rapidly progressive glomerulonephritis (RPGN).

Case details
A 21-year-old, newly married female without any living issue and foetal loss, resident of eastern India presented to IPGME&R hospital with presenting complaints of low to moderate intermittent fever (with occasional high rise of temperature) for last six months, pain, redness and swelling of both large and small joints (mostly additive) for last six months. Also rapidly progressive painful vesico-bullous skin lesion involving both sun exposed and non-exposed areas i.e. face, neck, upper back, limbs and
abdomen for last one month in multiple crops, with rupture of vesicles and draining of clear to turbid and occasionally haemorrhagic fluid. Some of the lesions subsided with scar formation. She also had history of scanty micturition for last 15 days with swelling of limbs and face for last 10 days.

Clinical examination revealed poor nutritional status with moderate pallor, bilateral pitting pedal oedema with facial puffiness, elevated jugular venous pressure (JVP) and raised temperature (101°F). Her pulse rate was 116/min, Blood pressure was 160/70 mm of Hg and respiration rate of 38/min.

Examination of skin revealed extensive vesicles and bullas with partial denudation in limbs, face, neck and in trunk. Nikolsky’s sign and bulla spread sign was positive. There was evidence of secondary infection and scarring (Figures 1-5).

Apart from inflammatory polyarthritis and left sided pleural effusion and mild hepato splenomegaly other systems were within normal limit.

Laboratory Examination shows:

- Blood Biochemistry and haematology: (Table 1),
- Chest X-RAY: Left more than right sided mild pleural effusion.
- Echocardiography: Mild Pericardial Effusion 10 cc, EF 65%, No features of Myocarditis.
- USG: Liver 15.9cm, normal shape and echotexture. Mild Splenomegaly with mild ascites. Kidney Right 11.8cm, left 11cm, CMD accentuated and bilateral cortical echogenicity increased, suggestive of renal parenchymal disease.
- Urine Routine: Proteinuria 3+, RBC 4-6/ HPF, Pus cell 1-2/HPF.
- Creatinine clearance: 20ml/min.
- 24 hour urinary protein estimation: 4.25gm in 24 hour. (Volume 1.45lt)
- Ascitic fluid study: Exudative with lymphocyte predominance.
- Pleural fluid study: Exudative, Low ADA, lymphocyte predominance.
- Serum ANA: ANA positive with 4+ homogeneous patterns with cytoplasmic position in 1:160 dilutions.
- ANA Profile: anti ds DNA 2+, anti-Ribosomal P protein 3+, anti-Histone +, anti-Smith +, anti SS-A +.
- C3 and C4: 28mg/dl and 6.24 mg/dl respectively.
- hs CRP: <0.03mg/dl. Ferritin: 771µgm/L.
- Thyroid and other hormonal profile were unremarkable.

- Skin Biopsy: Sub epidermal separation with neutrophilic infiltration in papillary dermis. Linear staining of the basement membrane zone with IgG, IgM, IgA, C3, and fibrinogen in direct immunofluorescence microscopy. Split Band is seen in the dermal side of the split (Floor pattern) with these Immuno reactants. Suggestive of Bullous lesion. Indirect immunofluorescence showed antibody to type VII collagen.
- Kidney Biopsy: Diffuse Lupus Nephritis Class IV of ISN/RPS classification with active lesion.

We diagnosed our patient as Bullous systemic lupus erythematosus (BSLE) with ISN/RPS class IV lupus nephritis with active lesion presenting with rapidly progressive glomerulonephritis (RPGN).

Initially patient was managed with Methylprednisolone pulse therapy 1gm once a day IV for three days followed by oral Prednisolone with Hydroxychloroquine and systemic broad spectrum antibiotics from day one. Dapsone was started with 100mg BD for bullous lesion but she was intolerant to Dapsone. Eventually her renal biopsy report came on day 10th of admission; with suggestion of active lupus nephritis, she was put on Rituximab 375mg/square meter body surface area. Later we received her skin biopsy report only to corroborate our clinical impression and forming our final diagnosis.

One month after completion of two doses of Rituximab, her renal impairment and skin lesions improved significantly. Biochemical assessment showed normalisation of creatinine value (1.2mg/dl) with a creatinine clearance of 60ml/min and reduction of proteinuria. In her subsequent visit after six months of 4th dose of Rituximab, proteinuria reduced further and renal recovery was nearly complete with creatinine clearance of 74ml/min without any further complication.

**Discussion**

BSLE accounts for less than one per cent of skin manifestations, generally affects young adult females and involves both sun-exposed and non-exposed areas. Because of the particularly clinical and histological presentation of BSLE, Camisa and Sharma proposed diagnostic criteria for BSLE; these include i) A diagnosis of SLE based on the ACR criteria; ii) Vesicles and bullae mainly located on sun-exposed areas; iii) The histopathology is characterised by sub-epidermal bullae with micro abscesses of neutrophils in the dermal papillae, similar to those found in dermatitis...
herpetiformis; and iv) Deposition of IgG, IgM, or both and often IgA in the basement membrane zone.\(^2\)

Rituximab which is a chimeric monoclonal antibody that reacts with CD20, an antigen that is present on immature, naive, and memory B cells but not on mature plasma cells, has been approved in the treatment of SLE with good efficacy and safety.\(^3\)

In this case report we have used Rituximab as an immuno modulator in a complicated life threatening situation of target organ damage. We have experienced a dramatic response in terms of skin lesions and renal recovery in our patient. Hydroxychloroquine and tapering dose of Prednisolone was prescribed to his patient and even after three successive follow up one month and six months apart no more bullous lesion was observed in this patient and her renal recovery was satisfactory.

Till now, there is only one case report showing improvement of skin lesion with Rituximab as a treatment of BSLE, but unlike this case Mycophenolate mofetil was used alongside.\(^4\) In this case a satisfactory response solely due to Rituximab has been observed even in an advanced life threatening nephropathy situation. This is obvious that potential role of Rituximab as an immuno modulator cannot be left aside.\(^5\)

**Conclusion**

Rituximab can be given as an alternative to Dapsone in bullous lesion associated with SLE and as an immuno modulator in active lupus nephritis with satisfactory renal outcome.

**References**


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**PEER REVIEW**

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**CONFLICTS OF INTEREST**

The authors declare that they have no competing interests.

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None

**PATIENT CONSENT**

The authors, Chakraborty A, Ghosh S, Ghosh S, Chowdhury A, Mondal D, declare that:

1. They have obtained written, informed consent for the publication of the details relating to the patient(s) in this report.
2. All possible steps have been taken to safeguard the identity of the patient(s).
3. This submission is compliant with the requirements of local research ethics committees.

**Figure 1:** Bullous lesion with secondary infection in neck
Figure 2: Bullous lesion with inflammatory hyper and hypopigmentation in face

Figure 3: Vesicobullous lesion

Figure 4: Skin biopsy showing Showing Bullous lesion with neutrophilic infiltrate and sub epidermal separation

Figure 5: Index patient after recovery

Table 1: Blood Biochemistry and Complete blood counts

<table>
<thead>
<tr>
<th>Complete Haemogram</th>
<th>Liver Function Test</th>
<th>Renal Function Test</th>
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<tbody>
<tr>
<td>Haemoglobin: 6.7gm/dl</td>
<td>Total Bilirubin:0.5mg/dl, AST:19U/L</td>
<td>Creatinine: 3.6mg/dl, Urea: 140mg/dl, Sodium: 128meq/lit.</td>
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<tr>
<td>TLC:7100/dl, N72, L24, E2</td>
<td>ALT:24U/L</td>
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<td>Platelets: 3.5×10^5/dl</td>
<td>Alkaline Phosphatase:38U/L</td>
<td>Potassium: 5.5meq/lit.</td>
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<tr>
<td>Microcytic Hypochromic peripheral blood picture, no abnormal cell.</td>
<td>Total Protein:5.4gm/dl</td>
<td>Calcium: 8.5mg/dl</td>
</tr>
<tr>
<td></td>
<td>Albumin:2.0gm/dl</td>
<td>Phosphate: 3.6mg/dl</td>
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<tr>
<td></td>
<td>Globulin:3.4gm/dl</td>
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