Efficacy of intravenous immunoglobulins in a patient with systemic lupus erythematosus presenting with Stevens-Johnson syndrome

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Accepted 11 July 2015

DESCRIPTION

A 38-year-old woman with systemic lupus erythematosus (SLE) manifested by tiredness, polyarthritis, lymphopaenia, elevated antidualle-stranded DNA antibodies, and hypocomplementaemia was initiated on azathioprine and hydroxychloroquine. After 2 weeks of treatment, she developed extensive erythematous raised skin patches, facial oedema, and desquamation of nasal and oral mucosa. Both drugs were discontinued and she improved within 4–7 days. Since hydroxychloroquine is associated with more hypersensitivity reactions than azathioprine, the latter was reintroduced, but 2 days later, the patient developed severe recurrent skin lesions (figure 1A–C). Skin biopsy showed band-like lymphocytic infiltrate with focal vacuolar interface changes consistent with early Stevens-Johnson syndrome.

Figure 1 Skin lesions on the (A) face, (B) abdomen and (C) back before intravenous immunoglobulin therapy.

Figure 2 Skin biopsy showing band-like lymphocytic infiltrate with focal vacuolar interface changes consistent with early Stevens-Johnson syndrome.
Our patient developed Stevens-Johnson syndrome most likely secondary to use of azathioprine. She had a remarkable clinical response to IVIG. Patients with SLE have an increased risk for Stevens-Johnson syndrome either because of the SLE itself or due to the use of immunosuppressive drugs such as azathioprine. Since cutaneous adverse events to azathioprine are rare, they are usually recognised on resumption of therapy. These reactions are dose dependent and occur within the first 4 weeks of exposure. There is no consensus on whether IVIG is better than conservative therapy for Stevens-Johnson syndrome, but some reports have shown its efficacy. In summary, we present an additional case in which high-dose IVIG was successful in accelerating the recovery period and stopping further disease progression of Stevens-Johnson syndrome.

Competing interests None declared.

Patient consent Obtained.

Provenance and peer review Not commissioned; externally peer reviewed.

REFERENCES

Learning points
- Patients with systemic lupus erythematosus are at increased risk of developing Stevens-Johnson syndrome.
- We described a 38-year-old woman who developed Stevens-Johnson syndrome secondary to azathioprine, and who had a rapid recovery with intravenous immunoglobulin treatment.
- This case together with other reports suggests that the use of intravenous immunoglobulins for Stevens-Johnson syndrome may be effective in accelerating the recovery period and stopping the progression of skin disease.