Pancytopenia as an early indicator for Stevens-Johnson syndrome complicated with hemophagocytic lymphohistiocytosis: a case report

Zhi-Dan Fan, Xiao-Qing Qian and Hai-Guo Yu*

Abstract

Background: Stevens-Johnson syndrome (SJS) is a severe skin and mucosal bullous disease. When complicated with Hemophagocytic lymphohistiocytosis (HLH), the condition is especially life-threatening.

Case presentation: Here we report the case of a 4-year-old boy suffering from SJS with extensive erythema multiforme and bulla. Despite active intervention and supportive care, the boy experienced increased skin lesions and a higher fever. Meanwhile, decreases in white blood cell count and hemoglobin were observed. Hyperferritinemia, increased soluble CD25 level, decreased NK cell activity and hemophagocytosis in the boy’s bone marrow confirmed the diagnosis of HLH. After high-dose intravenous immunoglobulin and methylprednisone pulse therapy, the boy was discharged in good condition.

Conclusion: Simultaneous occurrence of HLH and SJS is very uncommon and the condition is life-threatening. Pancytopenia can be a precocious indicator and enables to start a prompt diagnosis and treatment.

Keywords: Pancytopenia, Early diagnosis, Stevens-Johnson syndrome (SJS), Hemophagocytic lymphohistiocytosis (HLH)

Background

Stevens-Johnson syndrome (SJS) is characterized morphologically by the rapid onset of epidermal detachment and erosion of the mucous membrane. Epithelial loss and the subsequent bacterial and fungal infections are the most common cause of death in patients with SJS [1]. Immune dysregulation plays a key role in the pathogenesis of SJS. The death-inducing interaction of Fas with its Fas ligand triggers a suicidal caspase cascade in epidermal cells. High-dose intravenous immunoglobulin (IVIG) was shown to contain natural Fas-blocking antibodies and thus abrogate the keratinocyte apoptosis and the subsequent epidermal detachment [2].

Hemophagocytic lymphohistiocytosis (HLH), a lethal immune disorder, often leads to an abrupt onset of single organ failure and rapid progression to multiple organ failure [3]. Unlike familial HLH due to genetic defects leading to impaired functions of natural killer and cytotoxic T cells, secondary HLH is generally triggered by infections, immunosuppressant drugs, malignant or rheumatologic disorders [4]. So far, in patients with SJS, HLH has been rarely reported [5]. However, SJS complicated with HLH is a life-threatening condition as the diagnosis often occurs too late to start timely life-saving therapy. Here we report a 4-year-old boy who developed HLH following SJS. The laboratory tests of pancytopenia can be a precocious indicator for us to make a prompt diagnosis, and to start a timely treatment to ensure the successful outcome.

Case presentation

A previously healthy 4-year-old boy was admitted to our pediatric intensive care unit with a one-week history of spiking fever, coughing and a three-day history of severe mucous membrane and skin lesions. He received some cephalosporin and ibuprofen in another hospital for
acute upper respiratory tract infection. However, after four days of treatment, he developed a higher fever and a raised erythematous rash.

On admission, the boy appeared unwell with temperature above 39°C and heart rate of 192 beats/min. The examination showed oral mucositis, vesicobullous lesions and skin detachment across the body (Figure 1).

Routine blood tests were unremarkable on admission. Biochemistry parameters were normal except for albumin (27 g/L [normal 35–55]) and serum sodium (126 mmol/L [normal 135–145]). Both erythrocyte sedimentation rate (61 mm/h [normal < 20]) and C-reactive protein (85 mg/L [normal < 10]) were markedly increased. The pathological and bacteriologic examinations indicated the absence of viral, mycoplasma pneumoniae and chlamydia pneumoniae infections and negative results of blood culture and galactomannan test.

The active intervention, anti-infection measures, fluid compensation, electrolyte balance, nutritional support and intensive care were initiated immediately after hospitalization with SJS. Meanwhile, high-dose IVIG (2 g/kg, single continuous infusion) was given since IVIG can inhibit Fas-FasL interaction and halt the progression of SJS. HLH was suspected when the patient’s condition continued to deteriorate with hyperpyrexia and pancytopenia. Routine blood monitoring revealed anemia (hemoglobin 79 g/L), leukopenia (white blood cell count 0.6 x 10^9/L) and agranulocytosis (neutrophils 0.01 x 10^9/L). Then the HLH was suspected and the ferritin, NK cells and soluble CD25 were measured. Also the bone marrow smear was performed.

The diagnosis of SJS-associated HLH was established since the patient fulfilled 6 out of 8 HLH-2004 diagnostic criteria (Table 1) [4]. The laboratory tests showed hyperferritinemia (1031 μg/L), increased soluble CD25 level (8910 U/ml), decreased NK cell activity (1.02%) and hemophagocytosis in bone marrow (Figure 2).

Then pulse methylprednisolone (15 mg/kg/day) was administered for 3 consecutive days. The syndrome was considered to be in remission when the patient’s general condition improved, with normal temperature, ameliorated rash and normal routine blood examinations. Hereafter, methylprednisolone was continued at 2.5 mg/kg/day for another 3 days followed with appropriate rapid tapering by 20 mg every 3 days. His condition progressively improved, with normal hematological and biochemical laboratory parameters, inflammatory indices and ferritin level. On the day before discharge in good condition,
the bone marrow aspiration was performed showing the absence of hemophagocytosis. Some known genetic defects including PRF1, UNC13D and STX11 were examined to exclude the familial HLH but no mutations were found.

Discussion
Secondary HLH has been described mainly in association with viral, bacterial, fungal, and parasitic infections, malignant or rheumatologic disorders [4]. There is substantial mortality in secondary HLH, depending on the underlying disease [3]. So far, HLH has seldom been reported in patients with SJS. It should be noted that both SJS and HLH have a high mortality rate, so concurrence of the two in a patient is especially danger. Matsumoto Y. et al. [5] reported a 34-year-old woman who presented SJS complicated with HLH. Although repeated plasmapheresis was conducted, the woman still died of a brain hemorrhage due to disseminated intravascular coagulation.

Until recently, the pathophysiological mechanisms of secondary HLH are incompletely elucidated. The degree of cytokine disturbance is closely related to clinical signs and symptoms [6]. HLH often carries a very bad prognosis for being underdiagnosed and undertreated [7]. At the early stage before proinflammatory damage and cytokine storm, the opportunity to prompt recovery in effective treatment was often missed [8]. Furthermore, it is important for clinicians to differentiate HLH from underlying disease flares or infectious complications because they may share nearly all clinical and laboratory characteristics.

The timely awareness and diagnosis of HLH is the first, crucial step toward successfully life-saving therapy but is challenging because HLH is always complicated by relatively non-specific and variable presentations. NK-cell activity is typically low or absent in HLH, and should be also measured post remission [9]. CD25 is a useful inflammatory marker correlating with disease activity. But obtaining these two results required time. Hyperferritinemia is proved to be a valuable disease marker of HLH. An elegant study by Allen CE et al. [10] showed that a ferritin level over 10,000 ng/ml in children was 90% sensitive and 96% specific for HLH. Hemophagocytosis is important in the diagnostic work-up of HLH [4]. However, the sensitivity of hemophagocytosis was 83% with a specificity of only 60% for HLH [11]. Moreover, hemophagocytosis may be lacking early at onset. Waiting for evidence of hemophagocytosis to make therapeutic decisions in the bone marrow may delay appropriate therapy in this life-threatening condition [12]. It is important to note that these diagnostic criteria are mainly helpful in making the diagnosis instead of reflecting all of the typical clinical or laboratory features of patients with HLH. Of the ubiquitous responses in HLH, the pancytopenia is the most common feature and this can be easily detected by routine blood monitoring. HLH diagnosis is based on progressive cytopenia affecting at least two cell lineages. Considering the incidence of pancytopenia is very low in SJS, it can be an early indicator for SJS complicated with HLH and enables clinicians to start a prompt diagnosis and a timely life-saving therapy.

Therapeutic strategies for secondary HLH are not well established because randomized controlled clinical trials are not available [13]. However, based on our own experience [14] and some multi-center cohort studies [15], the initial treatment should be capable of suppressing the severe hyper-inflammation and controlling the underlying disease. A recent analysis of our experience treating 23 secondary HLH patients (so-called macrophage activation syndrome) triggered by systemic onset juvenile idiopathic arthritis, Kawasaki disease and systemic lupus erythematosus showed that 21 children were successfully treated with IVIG, corticosteroid and cyclosporine A. Corticosteroid and/or immunosuppressants seem to be the mainstay of treatment for secondary HLH [16]. When the trigger event is SJS per se in our patient, IVIG and corticosteroid are proposed to be first line therapeutic tools according to the HLH-2004 protocol [4]. High-dose IVIG can be given before a definite diagnosis of HLH since IVIG can halt the progression of SJS. In addition, we deemed that early use of IVIG may prevent a cytokine storm in HLH through the blockade of Fc receptors and other pathways. After establishing diagnosis of secondary HLH, the corticosteroid should be used to counteract the activated macrophages and stop the hyperinflammation stage of the syndrome.

Conclusions
In conclusion, HLH in SJS is very uncommon and the condition is life-threatening. IVIG in combination with corticosteroid is a safe and effective therapeutic regimen. Keeping highly vigilant to make an early diagnosis is essential to ensure favorable outcomes. Pancytopenia can be a precocious indicator and enables to start a prompt diagnosis and treatment.

Consent
Written informed consent was obtained from the patient’s parents for publication of this Case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

Abbreviations
SJS: Stevens-Johnson syndrome; HLH: Hemophagocytic lymphohistiocytosis; IVIG: Intravenous immunoglobulin; SHLH: Secondary hemophagocytic lymphohistiocytosis.

Competing interests
The authors declare that they have no competing interests.
We thank Lei Zhang for an invaluable help in revising our manuscript.

Acknowledgment

ZDF and XQQ drafted the manuscript. HGY made a critical revision. All authors read and approved the final manuscript.

References

1. Mockenhaupt M: The current understanding of Stevens-Johnson syndrome and toxic epidermal necrolysis. Expert Opin Immunother 2011, 12:803–813.
2. Chen J, Wang B, Zeng Y, Xu H: High-dose intravenous immunoglobulins in the treatment of Stevens-Johnson syndrome and toxic epidermal necrolysis in Chinese patients: a retrospective study of 82 cases. Eur J Dermatol 2010, 20:743–747.
3. Janka GE: Familial and acquired hemophagocytic lymphohistiocytosis. Eur J Pediatr 2007, 166:93–109.
4. Henter JI, Horne A, Aricó M, Egerer RM, Filipovich AH, Inashuki S, Ladisch S, McClain K, Webb D, Winiani J, Janka G: HLH-2004: diagnostic and therapeutic guidelines for hemophagocytic lymphohistiocytosis. Pediatr Blood Cancer 2007, 48:124–131.
5. Matsumoto Y, Naniwa D, Banno S, Sugura Y: The efficacy of therapeutic plasmapheresis for the treatment of fatal hemophagocytic syndrome: two case reports. Ther Apher 1998, 2:300–304.
6. Park HS, Kim DY, Lee JH, Lee JH, Kim SD, Park YH, Lee JS, Kim BY, Jeon M, Kang YA, Lee YS, Seol M, Lee YL, Lim YS, Jang S, Park CJ, Chi HS, Lee KH: Clinical features of adult patients with secondary hemophagocytic lymphohistiocytosis from causes other than lymphoma: an analysis of treatment outcome and prognostic factors. Ann Hematol 2012, 91:897–904.
7. Dhanankar M, Dassin SK: Hemophagocytic lymphohistiocytosis: a syndrome with diverse etiologies and treatment options. Clin Adv Hematol Oncol 2012, 10:262–265.
8. Larroche C: Hemophagocytic lymphohistiocytosis in adults: diagnosis and treatment. Joint Bone Spine 2012, 79:356–361.
9. Kardas F, Patiroglu T, Unal E, Chiang SC, Bryceson YT, Kendirci M: Hemophagocytic syndrome in a 4-month-old infant with biotinidase deficiency. Pediatr Blood Cancer 2012, 59:191–193.
10. Allen CE, Yu X, Kozinetz CA, McClain KL: Highly elevated ferritin levels and the diagnosis of hemophagocytic lymphohistiocytosis. Pediatr Blood Cancer 2008, 50:1227–1235.
11. Goel S, Potlik M, Imam H: Sensitivity and specificity of bone marrow hemophagocytosis in hemophagocytic lymphohistiocytosis. Ann Clin Lab Sci 2012, 42:1–25.
12. Gupta A, Tyrell P, Valari R, Benseler S, Weitzman S, Abdelhaleem M: The role of the initial bone marrow aspirate in the diagnosis of hemophagocytic lymphohistiocytosis. Pediatr Blood Cancer 2008, 51:402–404.
13. Demirkol D, Yildizdas D, Bayraktar B, Karapinar B, Kendirli T, Koroglu TF, Dursun O, Erkek N, Gedik H, Citak A, Kesici S, Karabucuoglu M, Carillo JA, Turkish Secondary HLH/MAS Critical Care Study Group: Hyperferritinemia in the critically ill child with secondary hemophagocytic lymphohistiocytosis/sepsis/multiple organ dysfunction syndrome/macrophage activation syndrome: what is the treatment? Crit Care 2012, 16:R52.
14. Ma HH, Qian XQ, Yu HG, Zhang YY, Fan ZD: Clinical analysis on 23 cases of childhood macrophage activation syndrome. J of Southeast Univ (Med Sci Edition) 2012, 31:736–738.
15. Takahashi N, Chubachi A, Kume M, Hatano Y, Komatsuda A, Kawaiata Y, Yanagiya N, Ichikawa Y, Miura AB, Miura I: A clinical analysis of 52 adult patients with hemophagocytic syndrome: the prognostic significance of the underlying diseases. Int J Hematol 2001, 74:209–213.
16. Lin CL, Yu HH, Lee JH, Wang LC, Lin YT, Yang YH, Chiang BL: Clinical analysis of macrophage activation syndrome in pediatric patients with autoimmune diseases. Clin Rheumatol 2012, 31:1223–1230.

Submit your next manuscript to BioMed Central and take full advantage of:

• Convenient online submission
• Thorough peer review
• No space constraints or color figure charges
• Immediate publication on acceptance
• Inclusion in PubMed, CAS, Scopus and Google Scholar
• Research which is freely available for redistribution

Cite this article as: Fan et al.: Pancytopenia as an early indicator for Stevens-Johnson syndrome complicated with hemophagocytic lymphohistiocytosis: a case report. BMC Pediatrics 2014 14:38.

doi:10.1186/1471-2431-14-38

www.biomedcentral.com/submit