Spontaneous Complete Remission in a Patient with Acute Myeloid Leukemia and Severe Sepsis

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Without treatment, acute myeloid leukemia (AML) is almost always fatal. Spontaneous remission of AML is a rare phenomenon and usually with a short duration [2]. It was first described following typhoid infection by Eisenlohr in 1878 [3,4].

Until 2014, according to modern criteria, only 46 eligible cases were found [5]. Several events like infection and blood transfusion are suggested to play an important role in SR. However, the exact mechanism of this phenomenon has yet to be determined [6]. One hypothesis is immune activation by these events has antileukemic effects [7]. Recent progression in early diagnosis and treatment makes SR infrequent [8].

We report a case of AML without cytogenetic abnormalities who had spontaneous complete remission after severe sepsis and both blood and platelet transfusion without chemotherapy.

1. Introduction

Acute myeloid leukemia (AML) manifests by proliferative, undifferentiated hematopoietic cells which infiltrate to blood, bone marrow, and other tissues. Approximately 10% of patient present with fever. Following chemotherapy, complete remission (CR) was defined with blood neutrophil and platelet count ≥1000/μL and ≥100000/μL, respectively, with no circulating blasts and blasts in bone marrow <5% [1]. Spontaneous remission (SR) without chemotherapy, complete or partial, is a very rare event with usually short duration [2]. It was first described following typhoid infection by Eisenlohr in 1878 [3,4].

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We report a case of AML without cytogenetic abnormalities who had spontaneous complete remission after severe sepsis and both blood and platelet transfusion without chemotherapy.

2. Case Report

A 53-year-old male presented with severe dyspnea, chest pain, productive cough, and dizziness. On admission, he was confused and pale with high-grade fever (39.5°C), tachypnea, and tachycardia, and blood pressure was 80/60 mmHg. There was no gingival hyperplasia and no peripheral lymphadenopathy or organomegaly. Chest X-ray revealed bilateral infiltrative abnormalities. He was transferred to Intensive care units (ICU) due to respiratory distress and mechanical ventilation started.

On admission, Laboratory findings were as follows: white blood cell (WBC), 400/μL, hemoglobin (Hgb), 6 g/dl, platelet (PLT) 54000/μL, C-reactive protein (CRP), 33 mg/L, and erythrocyte sedimentation rate (ESR), 125 mm/h. All of the renal and liver function tests, coagulation test, and electrolytes were within the normal values. The bone marrow aspiration smear (Figure 1) revealed 70%–80% cellularity with reduced megakaryocytes and raised ratio of myeloid to erythroid cells. Following bone marrow biopsy, the immunophenotyping analysis showed more than 80% blasts with positive CD13, CD14, CD33, CD34, CD43, CD64, HLA-DR, and c-MPO supported the AML-M4 based on French-American-British classification. Cytogenetic analysis demonstrated 46
Figure 1: Bone marrow smear at diagnosis showed 70%–80% cellularity with decreased megakaryocyte, increased proportion of myeloid to erythroid cells, and high level of myeloid blasts which could be presumptive of AML-M4.

Figure 2: Karyotype analyses showed 46 XY compatible with the apparently normal male with no chromosomal aberration on the basis of GTG technique at 350–400 band resolution.

XY compatible with the apparently normal male without chromosomal abnormality (Figure 2). The chemotherapy was postponed due to severe infection. The culture of tracheal discharge was positive for Enterobacter spp. and Acinetobacter baumannii. Blood culture result was Klebsiella pneumonia. Broad-spectrum antibiotics were started, and antifungal drug was added to antibiotics owing to persistent fever and neutropenia. The patient received red cells (10 units), platelet transfusions (10 units), and low-dose corticosteroid (dexamethasone 4 mg every 12 hr) during admission. Two weeks later, the infections completely resolved and he was weaned off the ventilator. Two weeks after sepsis resolution, hematologic status improved markedly with WBC of 3880/µL, Hgb of 9.2 g/dl, and PLT of 203000/µL. In addition, the bone marrow smear contained 5–10% cellularity with adequate megakaryocytes for this setting, myeloid and erytroid cells being in various maturation phases, and no blasts were recognized (Figure 3). Two weeks later, the bone marrow biopsy was repeated and the smear revealed 50–60% cellularity with sufficient megakaryocyte, the normal ratio of myeloid to erythroid cells, and no blast (Figure 4). The concurrent complete blood count (CBC) revealed WBC, 14600/µL, with neutrophil, 72.3%, Hgb, 9.4 g/dl, and PLT, 392000/µL (i.e., spontaneous complete remission). The immunophenotyping analysis showed normal plasma cell population. After 18 months, he has been in remission phase with WBC, 4800/µL, neutrophil, 69.0%, Hgb, 12.3 g/dl, and PLT, 253000/µL.

3. Discussion

According to the CR definition, our patient is in SCR until this manuscript is written (18 months). Among 46 eligible patients reported by Rashidi and Fisher as cases of SR, 39 patients achieved SCR with the median duration of remission only 5 months, whereas only nine patients had durable CR (defined as CR lasting for 1 year or longer). Therefore, our patient could be added to the latter group (Table 1) [9]. In other reports, the median period of remission was 8.2 ± 9.8 months [10].

Infections and blood transfusion lead to immune activation and have been frequently reported with the majority of SR cases like our case. However, exact mechanisms are not well known.

Moreover, cases unrelated to transfusions or infections have been documented previously. As in our case, pneumonia and bacteremia were significantly more common among CR cases compared to partial remission [5].
The author has no conflicts of interest.

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Table 1: Compare the characteristics of this case with previous similar cases reported by Rashidi and Fisher.

| Characteristics | Age | Sex | FAB | Karyotype | BM blasts (%) at presentation | Infection | Type | Fever | Irradiated transfusion | Remission | Duration | Relapse |
|----------------|-----|-----|-----|-----------|------------------------------|-----------|------|-------|----------------------|------------|---------|--------|
| This case      | 53  | M   | M4  | Normal    | >80%                         | +         | Pneumonia | +     | +                    | CR         | 18 m    | —      |
| Number of previously similar cases | 5   | 27  | 5   | 9         | 11                           | 32        | 13   | 37    | 32                   | 39         | 3       | 8      |

FAB: French-American-British classification; BM: bone marrow; CR: complete remission; m: months; M: male.