Acute respiratory distress syndrome associated with tumor lysis syndrome in a child with acute lymphoblastic leukemia

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Abstract

Tumor lysis syndrome is a serious and dangerous complication usually associated with antileukemia treatment in some malignancies characterized by high cell turn-over. Mild or severe electrolyte abnormalities including high serum levels of uric acid, potassium, phosphorus, creatinine, bun and reduction of calcium can be responsible for multi-organ failure, involving mostly kidneys, heart and central nervous system. Renal damage can be followed by acute renal failure, weight gain, progressive liver impairment, overproduction of cytokines, and subsequent maintenance of multi-organ damage. Life-threatening acute respiratory failure associated with tumor lysis syndrome is rare. We describe a child with T-cell acute lymphoblastic leukemia, who developed an unusually dramatic tumor lysis syndrome, after administration of the first low doses of steroid that was rapidly associated with severe acute respiratory distress syndrome (ARDS).

Case Report

A previously healthy 7-year-old boy with fever for seven days, headache, ocular pain and bilateral proptosis underwent a blood test that revealed marked leucocytosis, white blood cell count (WBC) 478,000/μL, slight anemia, red blood cells (RBC) 4,340,000/μL, hemoglobin (Hb) 10 g/dL, and mild thrombocytopenia (platelets 79,000/μL). He was immediately admitted to our Unit and a bone marrow aspirate confirmed the diagnosis of TLS. The patient was treated aggressively with intravenous fluids, allopurinol therapy, rasburicase, calcium gluconate, insulin therapy for hyperkalemia, and diuretics. Despite medical treatment, the patient was not able to maintain renal function parameters persistent. After administration of a total of 30 mg/kg of steroid in five doses in about a 36-hour-period, WBC dropped to 32,800/μL and he developed shock (pale and cold skin, small wrists, capillary Refill >4”, AP 78/57 mmHg, CF 132 bpm). Arterial blood gas value analysis revealed metabolic acidosis (pH 7.20, PO2 55 mmHg, PCO2 32 mmHg, HCO3- 12.5 mmol/L, BE -15.5, SatO2 79.8%). Blood tests showed uric acid 6 mg/dL, BUN 124 mg/dL, creatinine 2.9 mg/dL, potassium 6.9 mmol/L, calcium 4.3 mg/dL, phosphorus 26.6 mg/dL, supporting the diagnosis of TLS with acute renal failure.

The patient was treated aggressively with intravenous fluids, allopurinol therapy, rasburicase, calcium gluconate, insulin therapy for hyperkalemia, and diuretics. Due to anemia and thrombocytopenia he also needed red blood cell and platelet transfusions.

The following day he developed respiratory failure. A chest radiograph taken at this time revealed bilateral pulmonary infiltrates consistent with ARDS (Figure 1). Transfusion related acute lung injury (TRALI) was excluded because even though it is clinically indistinguishable from acute respiratory distress syndrome (ARDS), it occurs within 6 hours after transfusion. Unlike ARDS, TRALI is self-limiting, and there is usually clinical improvement within 48-96 hours if prompt respiratory support is provided.2 In our case significant levels of positive end-expiratory pressure and a high fraction of inspired oxygen (protective ventilation) were needed to maintain adequate systemic oxygenation. Despite medical treatment, hemofiltration (five sessions in three days) was necessary to control persistent hyper-
kalemia, hyperphosphemia with hypocalemia, and also to remove circulating inflammatory cytokines (probably related to ARDS). Steroids were continued at the dose of 10 mg/m²/day, in order to obtain an adequate control of leukemic cells count, and anti-inflammatory effect. He was intubated for one month and managed with continued mechanical ventilation. There was a gradual improvement of respiratory function, weaning from O2 therapy, and normalization of renal function with reversal of the altered hemodynamic factors. When the clinical condition improved he resumed chemotherapy according to the protocol, and currently he’s well in full complete remission, after discontinuation of chemotherapy, 36 months after diagnosis (laboratory findings are available in Supplementary Tables S1-S3).

Discussion

According to the consensus conference on the management of tumor lysis syndrome, laboratory TLS is defined by the occurrence of two or more of the following serum values before or after anticancer treatment (from three days before to seven days after the start of anticancer treatment): i) uric acid: increase of more than 25% from baseline, or 476 mmol/L (8 mg/dL); ii) potassium: increase of more than 25% from baseline, or 6.0 mmol/L (6 mEq/L); iii) phosphorus: increase of more than 25% from baseline, or 1.45 mmol/L (4.5 mg/dL) in adults, and 2.1 mmol/L (6.5 mg/dL) in children; iii) calcium: decrease of more than 25% from baseline, or 1.75 mmol/L (7 mg/dL).

Clinical TLS is defined by the presence of laboratory TLS and at least one of the following clinical alterations: renal failure (estimated glomerular filtration rate <60 mL/min), cardiac arrhythmia/sudden death, seizures. Grading of clinical TLS ranges from I to IV and is based on the degree of elevation of serum creatinine, the presence and type of cardiac arrhythmia, and the presence and severity of seizure.

Potential predictors of TLS include laboratory features, such as high initial WBC count, elevated serum LDH or uric acid levels, and clinical indicators of bulky disease, such as the presence of a mediastinal mass on chest radiographs, hepatomegaly (defined as a palpable liver 3 cm below the right costal margin), and splenomegaly (defined as a palpable spleen 2 cm below the left costal margin), as assessed by the physical examination on admission, pre-existing dehydration, oliguria, or renal failure and malignancies with high sensitivity to chemotherapy.

Drugs for TLS prophylaxis include allopurinol, which blocks the activity of the liver enzyme xanthine oxidase, and thereby decreases the risk of uric acid crystallization in the kidneys,^1^ and rasburicase, a recombinant urate oxidase enzyme, which converts existing uric acid to allantoin, which is 5 to 10 times more soluble in urine than uric acid. Effective treatment strategies that have been reported useful in TLS include vigorous hydration, allopurinol or rasburicase administration. Renal replacement therapy occasionally is needed. Indications for the start-up of renal replacement therapy in TLS include persistent hyperkalemia, severe metabolic acidosis, volume overload unresponsive to diuretic therapy, and overt uremic symptoms.

In our patient hypovolemic shock developed subsequently to TLS and acute renal failure; later severe ARDS began as a probable consequence of overproduction of cytokines. Despite medical treatment, continuous renal replacement therapy was necessary and after that only respiratory function gradually improved. To our knowledge, only one case of a 26-year-old woman with ARDS in TLS is reported in literature. The risk of developing ARDS in the setting of TLS may depend on the degree of cell lysis, the speed of release of mediators into pulmonary circulation, and the susceptibility of the alveolar-capillary membrane to injury.

The pathogenesis of ARDS in the setting of TLS remains unclear. It could be related to the release of cytokines or other active mediators, and not to the serum electrolyte abnormalities. It has been hypothesized that an excessive inflammatory response could be down-regulated with continuous renal replacement therapies (CRRT), by non specific extracorporeal removal of cytokines, and other mediators. Absorption appears to be the predominant mechanism of mediator elimination. In addition, the eventual elimination of inflammatory mediators, fluid removal with reduction of extravascular lung water (EVLW), is a second mechanism by which hemofiltration may be beneficial in ARDS. However, the observed hemodynamic improvement can at least partially be attributed to a reduction of body temperature, and the evidence for clinically important removal of pro-inflammatory cytokines remains limited. CRRT-induced hypothermia can be used in patients with ARDS in order to reduce CO2 production. The decreased ventilator requirement reduces the risk of ventilator-induced lung injury. Reduced CO2 production combined with the alkalinizing effect of bicarbonate in the replacement solution facilitates the institution of permissive hypercapnia. An advantage of this procedure is the ability to remove large volumes of fluid while avoiding the hypotensive episodes caused by intermittent hemodialysis, so it is indicated for managing patients with acute renal failure who are hemodynamically unstable.

Figure 1. Chest radiograph shows multiple bilateral pulmonary infiltrates suggesting an acute respiratory distress syndrome.

Conclusions

Tumor lysis syndrome (TLS) is a very serious and potentially life-threatening event, that can lead to severe multi-organ dysfunction. Intensive supportive treatment, including hyperhydration and allopurinol could be not enough in some malignancies characterized by a rapid cellular turn over and the administration of rasburicase represent the first choice prophylaxis, that must be started as soon as possible. ARDS can be an unusual complication of TLS, probably related to overproduction of cytokines, also in spite of steroid treatment. A prompt hemofiltration may be an effective measure for both clinical conditions, clearing the excess of circulating cytokines and supplying renal function.

References

5. Sayah DM, Looney MR, Toy P. Transfusion reactions: newer concepts on the pathophysiology, incidence, treatment, and pre-

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