Acute Tumor Lysis Syndrome during Oral Fludarabine Treatment for CLL – a Rare Event that Might Be Observed More Frequently in the Future

Catherine Rioufol, Bertrand Coiffier

Pharmacy, Centre Hospitalier Lyon-Sud, Department of Hematology, Centre Hospitalier Lyon-Sud and Université Claude Bernard, Lyon, France

Tumor lysis syndrome (TLS) with acute renal failure is a very rare complication of oral fludarabine therapy in patients with chronic lymphocytic leukemia (CLL). Cheson reviewed 6,137 patients with CLL treated with fludarabine and found an incidence of TLS in 0.33% of the patients [2]. TLS has also been observed in CLL patients treated with oblimersen, flavopiridol, high-dose corticosteroids, and rituximab [3–6].

TLS is caused by the lysis of malignant cells by cytotoxic chemotherapy, leading to a rapid release of intracellular contents into the blood stream including potassium, phosphate, and purines from the breakdown of nuclear proteins. Secondary hypocalcemia, hyperkalemia and hyperphosphatemia lead to gastrointestinal disturbances, neuromuscular effects and cardiovascular consequences. Purines are catabolized to hypoxanthine, then xanthine, and finally in the liver to uric acid by the enzyme xanthine oxidase. The precipitation of excess uric acid contributes to impaired renal function; acute renal failure represents the most frequent and serious clinical consequence of TLS-related hyperuricemia [7]. Risk factors include the type of malignancy, some tumor-related factors, such as high numbers of circulating tumor cells and a large tumor burden, the presence of individual factors, such as pre-existing renal insufficiency, and the type and intensity of anticancer regimen used [7].

A literature survey in 2003 reported 5 cases of intravenous fludarabine-induced TLS in patients with CLL that occurred within the first week of initiation of therapy [8]. However, the 20 cases reported by Cheson in 1998 were not accounted for in this survey [2]. Since then some other case reports have been published but they are rare because most of the patients at risk received allopurinol prophylaxis. It is probable that the incidence is higher, most cases not being reported anymore.

Since allopurinol prophylaxis is available for oral use in Europe, only one case of oral fludarabine-induced TLS was described [9]. The patient, which had been treated with oral fludarabine 40 mg/m² daily for 5 days, developed 2 episodes of acute TLS 17 and 15 days after starting fludarabine for the...
first and second treatment. The authors concluded that fludarabine appeared to cause TLS later after oral administration compared to intravenous therapy. In the case reported in this issue, TLS occurred at day 7.

The standard management strategy for treating TLS is based on hydration and decreasing uric acid concentration. This may be obtained with rasburicase, a newly developed recombinant urate oxidase, that converts uric acid to allantoin, a much more soluble product [10, 11]. Several studies have demonstrated that rasburicase is generally well-tolerated, although severe adverse effects were reported such as hypersensitivity reactions [10]. Allopurinol has a great efficacy for preventing TLS in high-risk patients but has a limited activity when TLS is installed and creatinine levels have increased.

It has been suggested that CLL patients should receive allopurinol routinely during the first course of fludarabine therapy; however, we think that it has only a role in high-risk patients [10]. These patients may be defined by a high blood lymphocyte count (greater than $10^6$ G/l) or high tumor mass (multiple lymph nodes larger than 5 cm). In a high-risk patient allopurinol should be given for 10 days because TLS is often delayed with fludarabine. In case of oral fludarabine, 15–20 days might be better because 2 of the 3 reported incidences occurred after 15 days.

Calvo-Villas’s paper confirms the strong clinical benefits of the newly developed recombinant urate oxidase rasburicase, to achieve a complete recovery of renal function and a decrease of the uric acid level to within the normal range in patients with acute renal failure secondary to TLS. The short 2-days treatment was sufficient but sometimes 3–7 days are necessary to normalize uric acid and creatinine levels. In the case of CLL and fludarabine where TLS occurs later than with chemotherapy in lymphomas, a short treatment is usually sufficient [12]. Furthermore, patients themselves should be educated about the symptoms associated with TLS and the need for high fluid intake during the first course of treatment.

In conclusion, TLS is a rare event in CLL patients but one that may occur more frequently because of the very active armamentarium that is currently developed for these patients. It usually occurs 7–10 days after treatment. It is easily prevented with allopurinol and must be treated with hydration plus rasburicase.

References