

# ONCOLOGY EMERGENCY ESSENTIALS: Addressing Tumor Lysis Syndrome in Your Practice

## Risk Factors



- Malignancies with large, rapidly growing cells (bulky tumors; high proliferation rate)
- Tumors that are chemosensitive; novel/effective cancer treatments
- High tumor burden (↑ stage, WBC counts, LDH levels)
- Patients with pre-existing renal dysfunction/involvement

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Measure serum uric acid, potassium, phosphate, and calcium  
If  $\leq 1$  abnormal value

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TLS Risk Stratification According to Malignancy

LOW RISK	INTERMEDIATE RISK	HIGH RISK
Most solid tumors	Bulky or advanced solid tumors sensitive to chemotherapy, such as neuroblastomas, germ-cell tumors, and small-cell lung cancer	
Myelomas		
CML	CLL treated with targeted and/or biological therapies and/or those with $WBC \geq 50 \times 10^9/L$	
CLL treated with alkylating agents		
AML with $WBC < 25 \times 10^9/L$ and $LDH < 2 \times ULN$	AML with $WBC < 25 \times 10^9/L$ and $LDH \geq 2 \times ULN$ , or $WBC 25-100 \times 10^9/L$	AML with $WBC \geq 100 \times 10^9/L$
	ALL with $WBC < 100 \times 10^9/L$ and $LDH < 2 \times ULN$	ALL with $WBC < 100 \times 10^9/L$ and $LDH \geq 2 \times ULN$ , or $WBC \geq 100 \times 10^9/L$
		B-ALL
Hodgkin lymphoma Indolent NHL	Stage I/II BL and LBL with $LDH < 2 \times ULN$	Stage III/IV BL and LBL, or Stage I/II with $LDH \geq 2 \times ULN$
Adult ALCL; Child stage I/II ALCL	Child stage III/IV ALCL	
Adult ATL, DLBCL, PTCL, transformed, and mantle cell (blastoid variants) lymphomas with $LDH \leq WNL$ ; Child stage I/II	Adult ATL, DLBCL, PTCL, transformed, and mantle cell (blastoid variants) lymphomas with $LDH > ULN$ and non-bulky; Child stage III/IV with $LDH < 2 \times ULN$	Adult ATL, DLBCL, PTCL, transformed, and mantle cell (blastoid variants) lymphomas with $LDH > ULN$ and bulky; Child stage III/IV with $LDH \geq 2 \times ULN$

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Renal Function Adjustment

Low-risk lymphoma/leukemia with **renal dysfunction/involvement**

Intermediate-risk disease with **renal dysfunction/involvement** or with uric acid, potassium, and/or phosphate  $> ULN$

ALCL, anaplastic large cell lymphoma; ALL, acute lymphocytic leukemia; AML, acute myeloid leukemia; ATL, adult T-cell lymphoma; BL, Burkitt lymphoma/leukemia; CLL, chronic lymphocytic leukemia; CML, chronic myeloid leukemia; DLBCL, diffuse large B-cell lymphoma; LBL, lymphoblastic lymphoma; LDH, lactate dehydrogenase; NHL, non-Hodgkin lymphoma; PTCL, peripheral T-cell lymphoma; ULN, upper limit of normal; WBC, white blood cells; WNL, within normal limits.

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If  $\geq 2$  abnormal values of serum uric acid, potassium, phosphate, or calcium

Element	Value
Uric acid	$\geq 476 \mu\text{mol/L}$ (8.0 mg/dL)
Potassium	$\geq 6.0 \text{ mmol/L}$ (6.0 mEq/L)
Phosphorous	$\geq 2.1 \text{ mmol/L}$ (6.5 mg/dL) for children $\geq 1.45 \text{ mmol/L}$ (4.5 mg/dL) for adults
Calcium	$\leq 1.75 \text{ mmol/L}$ (7.0 mg/dL)

+  $\geq 1$  of acute kidney injury, cardiac dysrhythmia, or seizure

Laboratory TLS

Clinical TLS

## Prophylaxis Recommendations

LOW RISK

INTERMEDIATE RISK

HIGH RISK

Monitor TLS parameters, including uric acid, potassium, phosphate, calcium, creatinine, and LDH, as well as urine output

+ Normal hydration

+ Increased hydration  
(3L/m<sup>2</sup>/day)

+ Increased hydration  
(3L/m<sup>2</sup>/day)

No prophylaxis for hyperuricemia

+ Allopurinol  
100–300 mg, po, q8h, daily  
(can consider rasburicase)

+ Rasburicase  
one dose, 0.1–0.2 mg/kg IV

## Management of Established Clinical TLS

Admission to intensive care unit

Cardiac monitoring

Monitoring TLS laboratory abnormalities every 2–4 hours

+ Vigorous hydration

+ Correction of electrolyte abnormalities  
(management of hyperkalemia and/or hyperphosphatemia)

+ Rasburicase

### References:

Coiffier B, Altman A, Pui CH, Younes A, Cairo MS. Guidelines for the management of pediatric and adult tumor lysis syndrome: an evidence-based review. *J Clin Oncol.* 2008;26(16):2767-2778.

Cairo MS, Coiffier B, Reiter A, Younes A, TLS Expert Panel. Recommendations for the evaluation of risk and prophylaxis of TLS in adults and children with malignant diseases: an expert TLS panel consensus. *Br J Haematol.* 2010;149(4):578-586.