Rasburicase Dosing Criteria

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<u>Background:</u> Rasburicase (Elitek) is indicated for initial management of plasma uric acid levels in pediatric and adult patients with leukemia, lymphoma, and solid tumor malignancies who are receiving anti-cancer therapy expected to result in tumor lysis and subsequent elevation of plasma uric acid. Approved dosing is at 0.2 mg/kg as an intravenous infusion over 30 minutes daily for up to 5 days, but fixed dosing strategies have also been described.¹⁻⁷

Based on a Rasburicase MUE presented at the February 2015 P&T meeting, the following criteria are proposed:

- 1. Flat dosing: Adult patients would receive a maximum of rasburicase 6 mg IV X 1 dose
- 2. Rasburicase criteria for use:

To receive a dose of rasburicase, patients should <u>one</u> have the following criteria:

- Established diagnosis of malignancy associated with a high risk of tumor lysis syndrome (TLS) such as:
 - Acute leukemias with WBC >50,000
 - Burkitt's or lymphoblastic lymphoma
 - Other lymphomas with bulky disease

Or <u>two</u> of the following:

- Hematologic malignancy which may be associated with TLS but not listed above
- Solid tumors with high proliferative rates and rapid response to therapy (germ cell tumor, small cell lung cancer, etc.)
- Uric acid >7.5 mg/dL
- Signs of renal dysfunction in patients with a malignancy diagnosis
 - SCr >1.5XULN (1.875) or oliguria
 - Patients that are scheduled to receive dialysis should not be considered
- 3. Patients may receive an additional dose of rasburicase 6 mg IV X 1 within 7 days if the uric acid >2.5 mg/dL.
- 4. Laboratory: Uric acid levels obtained for 4 days after a dose of rasburicase will be ordered under a separate Cerner orderable. Rasburicase and the uric acid levels will be ordered as part of a Powerplan.

References:

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- 9. Coiffier B, Altman A, Pui CH, et al. Guidelines for the management of pediatric and adult tumor lysis syndrome: an evidence-based review. *J Clin Oncol*. 2008; 26:2767-78.
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