BC Cancer Protocol Summary for Therapy for Indolent Lymphoma and Chronic Lymphocytic Leukemia Using Chlorambucil

Protocol Code LYCHLOR

Tumour Group Lymphoma

Contact Physician Dr. Laurie Sehn

ELIGIBILITY:

- Malignant lymphoma, indolent, including follicular, lymphoplasmacytic, small lymphocytic and marginal zone lymphomas
- Chronic lymphocytic leukemia

EXCLUSIONS:

Active hemolytic anemia or immune-related thrombocytopenia

TESTS:

- Baseline (required before first treatment): CBC & diff, platelets, bilirubin, ALT
- Baseline (required, but results do not have to be available to proceed with first treatment; results must be checked before proceeding with further treatment): HBsAg, HBcoreAb
- Before each treatment: CBC & differential, platelets

PREMEDICATIONS:

None

SUPPORTIVE MEDICATIONS:

If HBsAg or HBcoreAb positive, start lamiVUDine 100 mg PO daily for the duration of chemotherapy and continue for one year from treatment completion for patients who are HBsAg positive and for six months for patients who are HBcoreAb positive.

TREATMENT:

Three available schedules, choice determined by individual patient characteristics.

Drug	Dose	BC Cancer Administration Guideline
Schedule 1: chlorambucil	Starting dose: 0.4 mg/kg for one dose on day 1 every 2 weeks. Subsequently, if ANC greater than 3.5 x 10 ⁹ /L, increase dose by 0.1 mg/kg, adjusting dose to induce a therapeutic response but not cause a fall in neutrophil count below 1.2 x 10 ⁹ /L. MAXIMUM DOSE: 0.8 mg/kg every 2 weeks.	PO
	Round dose to the nearest 2 mg. Administer on an empty stomach.	
OR		
Schedule 2: chlorambucil	0.2 mg/kg once daily for 21 consecutive days (total dose per cycle 4.2 mg/kg) adjusted to induce a therapeutic response but not cause a fall in neutrophil count below 1.2 x 10 ⁹ /L. Repeat every 6 weeks.	PO
	Round dose to the nearest 2 mg. Administer on an empty stomach.	
OR		
Schedule 3: chlorambucil	0.1mg/kg once daily (range 0.03-0.2 mg/kg once daily) continuously adjusted to induce a therapeutic response but not cause a fall in neutrophil count below 1.2 x 10 ⁹ /L.	PO
	Round dose to the nearest 2 mg. Administer on an empty stomach.	

Continue treatment until two months <u>after</u> maximum response achieved (maximum 1 year)

DOSE MODIFICATIONS:

Hematological, for low counts due to treatment, not disease

ANC (x10 ⁹ /L)	Platelets (x10 ⁹ /L)	Dose (all drugs)
Greater than or equal to 1.2	Greater than or equal to 80	100%
Less than 1.2	Less than 80	Delay until recovery

PRECAUTIONS:

- 1. **Neutropenia**: Fever or other evidence of infection must be assessed promptly and treated aggressively.
- 2. Hepatitis B Reactivation: All lymphoma patients should be tested for both HBsAg and HBcoreAb. If either test is positive, such patients should be treated with lamiVUDine during chemotherapy and continue for one year from treatment completion for patients who are HBsAg positive and for six months for patients who are HBcoreAb positive. Such patients should also be monitored with frequent liver function tests and hepatitis B virus DNA at least every two months. If the hepatitis B virus DNA level rises during this monitoring, management should be reviewed with an appropriate specialist with experience managing hepatitis and consideration given to halting chemotherapy.

Call Dr. Laurie Sehn or tumour group delegate at (604) 877-6000 or 1-800-663-3333 with any problems or questions regarding this treatment program.