

## Chronic Lymphocytic Leukaemia (CLL): Recommendations for GP Monitoring

### Presentation

- Your patient has been diagnosed with CLL.
- This is the most common haematological malignancy in the western world with 1 in 200 patients developing it at some point in their life.
- It is a blood and bone marrow condition that usually progresses slowly over a number of years with the accumulation of these abnormal cells in the bone marrow and lymph glands.
- It is this accumulation of cells which can often be detected by changes in the person's blood or on physical examination.
- About 10-25% patient with CLL never need any treatment but do need to be monitored for any change.

### Clinical Findings

Your patient does not require treatment at present. This will usually be because your patient has Binet stage A CLL (see below). We would be grateful if you could monitor every 6 months with a full blood count together with a clinical assessment for signs and symptoms that could be due to progressive CLL. These include:

- Progressive lymphadenopathy or splenomegaly
- Systemic symptoms of weight loss, drenching sweats, unexplained fevers or fatigue

### Binet staging system

Binet staging system is used to describe the stage of the CLL. Patients with Stage A disease can often be safely managed in general practice. Patients with stage B and C disease will usually be actively monitored or treated at hospital.

**Stage A** – fewer than 3 groups of enlarged lymph nodes detected with no anaemia or thrombocytopenia.

**Stage B** – More than 3 groups of enlarged lymph nodes with no anaemia or thrombocytopenia.

**Stage C** – Enlarged lymph nodes or spleen and anaemia or thrombocytopenia.

## Investigations

6 monthly investigations	Investigations to consider
FBC	Infection screen (if fever / new sweats)
U+E / LFTs	Lactate dehydrogenase
History and examination	Immunoglobulins (if recurrent infection)
	Haematinics (if new anaemia)
	Haemolysis screen: please discuss if concerns about associated haemolytic anaemia.

## Management

### Vaccinations

Patients with CLL should receive the annual flu vaccination and should be kept up to date with vaccinations for pneumococcus, haemophilus and meningococcus with repeats every 5 years. Please withhold vaccinations post treatment until lymphocyte count is  $> 1$  or  $> 6$  months post chemotherapy.

**Please do not give these patients any live vaccines including the shingles vaccine.**

Patients with CLL can be prone to getting chest infections due to underlying immunoparesis with some patients developing bronchiectasis. These patients should be managed with the haematologist and respiratory teams.

### Referral

Indications for referral back to hospital include

- Progressive fall in patients' blood counts (without other cause): Hb  $< 100$  g/L or Platelets  $< 100 \times 10^9$ /L.
- A lymphocyte count of  $>100 \times 10^9$ /L
- A lymphocyte count that doubles over a 6 month period once the lymphocytes are  $>30 \times 10^9$ /L
- New lymphadenopathy in  $> 3$  regions
- Rapidly enlarging lymph nodes
- Persistent symptoms of fatigue/anorexia/weight loss ( $>10\%$  in 6 months)
- Drenching night sweats (changing linen  $> 2$  times a week for  $>2$  weeks), recurrent infections or unexplained fevers.

### References