

Lymphoid Conditions Clinical Management Protocol

Protocol	Eligibility		Monitoring	Recall
<p>Chronic Lymphocytic Leukaemia (CLL) & Small Lymphocytic Leukaemia (SLL)</p> <p>LEVEL 2 PIFU</p> <p><i>Review at Year 25</i></p>	<p>No ongoing functional or psychological problems requiring secondary care input identified on the HNA & fulfil one of the 3 scenarios below (all criteria must be met):</p> <ul style="list-style-type: none"> • Consultant decision to recruit to PIFU. • Not participating in a clinical interventional trial. • Recruited in MLMS study (at consultant discretion) <p>One of the following scenarios:</p> <p>Scenario 1 –</p> <ol style="list-style-type: none"> Low-risk patient according to Consultant criteria [including Untreated Rai stage 0 or clinical MBL, Mutated IGHV, No evidence of high-risk molecular lesions (11q deletion, Trisomy 12, 17p deletion, TP53 mutation)] No progressive or active CLL No complications or concomitant conditions requiring haematologist intervention (e.g. recurrent infections, chronic cytopenia's) <p>Scenario 2 –</p> <ol style="list-style-type: none"> Following finite treatments (e.g., Ibrutinib+venetoclax, venetoclax+obinutuzumab or venetoclax+rituximab) and EOT assessment of response indicates radiological CR (by CT and or PET/CT), normal blood counts and undetectable MRD in the peripheral blood using HMDS criteria. No treatment-related toxicity requiring haematologist intervention. <p>Scenario 3 –</p> <ol style="list-style-type: none"> Following any treatments Low-risk recurrence according to Consultant criteria [including Mutated IGHV, No evidence of high-risk molecular lesions (11q deletion, Trisomy 12, 17p deletion, TP53 mutation)] EOT assessment of response indicates radiological CR (by CT and or PET/CT), normal blood counts, no treatment-related toxicity requiring haematologist intervention. No evidence of active or progressive disease. <p>Scenario 4 –</p> <ol style="list-style-type: none"> high-risk patient No progressive or active CLL for more than 5 years No complications or concomitant conditions requiring haematologist intervention (e.g. recurrent infections, chronic cytopenia's) 	<p>Exclude patients who cannot self-manage or must attend the clinic to manage functional or psychological issues.</p>	<p>Year 1 – Month 6 & 12</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 2 – Month 18 & 24</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 3 – Month 30 & 36</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 4 – 25 Annual Surveillance –</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire 	<p>Trend changes/new abnormalities in blood results</p> <ul style="list-style-type: none"> • Hb <110 • Platelets <100 • Neutrophils <1.0 • Lymphocyte count doubles over 30 in <6 months <p>Patient reports signs or symptoms lasting more than 4 weeks & not explained by other conditions -</p> <ul style="list-style-type: none"> • New enlarged lymph nodes • Night sweats • Recurrent infections in the last 12months (>3) • Abdominal discomfort, feeling full early when eating or after eating a small amount of food • Shortness of breath at rest or on exertion • Worsening tiredness/fatigue • Unexplained weight loss or loss of appetite • Changes in bowel habits

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<p>Leukaemic MZL, including Splenic MZL</p> <p>LEVEL 2 PIFU</p> <p><i>Review at Year 25</i></p>	<p>No ongoing functional or psychological problems requiring secondary care input identified on the HNA & fulfil one of the 2 scenarios below (all criteria must be met):</p> <ul style="list-style-type: none"> • Consultant's decision to recruit to PIFU. • Not participating in a clinical interventional trial. • Recruited in MLMS study (at consultant discretion) <p>One of the following scenarios:</p> <p>Scenario 1 –</p> <ol style="list-style-type: none"> post diagnosis if untreated Low-risk patient according to Consultant criteria <p>Scenario 2 –</p> <ol style="list-style-type: none"> Post completion of treatment Complete, or stable partial remission confirmed on: <ul style="list-style-type: none"> • CT scan or clinical examination (at the Consultant's discretion) • Blood results 	<p style="writing-mode: vertical-rl; transform: rotate(180deg);">Exclude patients who cannot self-manage or must attend the clinic to manage functional or psychological issues.</p>	<p>Year 1 – Month 6 & 12</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 2 – Month 18 & 24</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 3 – Month 30 & 36</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 4 – 25 Annual Surveillance –</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire 	<p>Trend changes/new abnormalities in blood results</p> <ul style="list-style-type: none"> • Hb <110 • Platelets <100 • Neutrophils <1.0 • Lymphocyte count doubles over 30 in <6 months <p>Patient reports signs or symptoms lasting more than 4 weeks & not explained by other conditions -</p> <ul style="list-style-type: none"> • New enlarged lymph nodes • Night sweats • Recurrent infections in the last 12months (>3) • Abdominal discomfort, feeling full early when eating or after eating a small amount of food • Shortness of breath at rest or on exertion • Worsening tiredness/fatigue • Unexplained weight loss or loss of appetite • Changes in bowel habits

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<p>Hairy Cell Leukaemia (HCL) & Variant Hairy Cell Leukaemia (HCLv)</p> <p>LEVEL 2 PIFU</p> <p><i>Review at Year 25</i></p>	<p>No ongoing functional or psychological problems requiring secondary care input identified on the HNA & fulfil one of the 2 scenarios below (all criteria must be met):</p> <ul style="list-style-type: none"> • Consultant's decision to recruit to PIFU. • Not participating in a clinical interventional trial. • Recruited in MLMS study (at consultant discretion) <p>One of the following scenarios:</p> <p>Scenario 1 –</p> <ol style="list-style-type: none"> a. post diagnosis if untreated b. Low-risk patient according to Consultant criteria <p>Scenario 2 –</p> <ol style="list-style-type: none"> a. Post completion of treatment b. Complete, or stable partial remission confirmed on: <ul style="list-style-type: none"> • CT scan or clinical examination (at the Consultant's discretion) • Blood results 	<p>Exclude patients who cannot self-manage or must attend the clinic to manage functional or psychological issues.</p>	<p>Year 1 – Month 6 & 12</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 2 – Month 18 & 24</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 3 – Month 30 & 36</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 4 – 25 Annual Surveillance –</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire 	<p>Trend changes/new abnormalities in blood results</p> <ul style="list-style-type: none"> • Hb <110 • Platelets <100 • Neutrophils <1.0 • Lymphocyte count doubles over 30 in <6 months <p>Patient reports signs or symptoms lasting more than 4 weeks & not explained by other conditions -</p> <ul style="list-style-type: none"> • New enlarged lymph nodes • Night sweats • Recurrent infections in the last 12months (>3) • Abdominal discomfort, feeling full early when eating or after eating a small amount of food • Shortness of breath at rest or on exertion • Worsening tiredness/fatigue • Unexplained weight loss or loss of appetite • Changes in bowel habits

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<p>Leukaemic T-cell lymphomas</p> <p>LEVEL 2 PIFU</p> <p><i>Review at Year 25</i></p>	<p>No ongoing functional or psychological problems requiring secondary care input identified on the HNA & fulfil one of the 2 scenarios below (all criteria must be met):</p> <ul style="list-style-type: none"> • Consultant's decision to recruit to PIFU. • Not participating in a clinical interventional trial. • Recruited in MLMS study (at consultant discretion) <p>One of the following scenarios:</p> <p>Scenario 1 –</p> <ol style="list-style-type: none"> post diagnosis if untreated Low-risk patient according to Consultant criteria <p>Scenario 2 –</p> <ol style="list-style-type: none"> Post completion of treatment Complete, or stable partial remission confirmed on: <ul style="list-style-type: none"> • CT scan or clinical examination (at the Consultant's discretion) • Blood results 	<p style="writing-mode: vertical-rl; transform: rotate(180deg);">Exclude patients who cannot self-manage or must attend the clinic to manage functional or psychological issues.</p>	<p>Year 1 – Month 6 & 12</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 2 – Month 18 & 24</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 3 – Month 30 & 36</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire <p>Year 4 – 25 Annual Surveillance –</p> <ul style="list-style-type: none"> • FBC • Offered a Symptom Questionnaire 	<p>Trend changes/new abnormalities in blood results</p> <ul style="list-style-type: none"> • Hb <110 • Platelets <100 • Neutrophils <1.0 • Lymphocyte count doubles over 30 in <6 months <p>Patient reports signs or symptoms lasting more than 4 weeks & not explained by other conditions -</p> <ul style="list-style-type: none"> • New enlarged lymph nodes • Night sweats • Recurrent infections in the last 12months (>3) • Abdominal discomfort, feeling full early when eating or after eating a small amount of food • Shortness of breath at rest or on exertion • Worsening tiredness/fatigue • Unexplained weight loss or loss of appetite • Changes in bowel habits