

HAEMATOLOGY GP Pathway Guides

V.4 March2026

Please note: these guidelines are specific to MFT. While local guidelines are similar across Manchester, please refer to local policy

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16. Hyperferritinaemia
17. Easy bruising

Epidemiological Notice for Greater Manchester: As Greater Manchester is a high HIV prevalence area, clinicians are strongly encouraged to consider routine HIV testing for all adult patients presenting with unexplained haematological or lymphoproliferative features, regardless of perceived individual risk profiles.

B Symptoms

- Weight loss >10% over 6 months
- Drenching sweats,
- Unexplained fever >38°C

Lymphadenopathy

Lymphadenopathy- look for causes

Lymphadenopathy associated with:

- B symptoms
- Liver and spleen enlargement
- Rapidly increasing in size
- Generalised lymphadenopathy
- Cytopenias

Refer to Haematology on urgent (suspected cancer) pathway

Localised unexplained adenopathy
OR
Concerns of metastatic node

Appropriate referral to surgical team or ENT for biopsy/ radiological biopsy (US or CT guidance)

Causes:

- Acute and chronic bacterial infections
- Syphilis
- Auto immune conditions
- Malignancy (haematological/ metastatic)
- Viral infections (including HIV, EBV, CMV)

Patient Information
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Referral Proforma
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B symptoms

- Weight loss >10% over 6 months
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Splenomegaly

Spleen >14cm

NB: Spleen increases with height. On average increases in length by 0.2 cm for every inch > 6ft

Consider using expected spleen size calculator by QxMD
Calculate:

https://qxmd.com/calculate/calculator_384/expected-spleen-size

- B symptoms
- Cytopenias
- Increased LDH
- Paraprotein
- Lymphadenopathy
- High haemoglobin or increased platelet count
- Evidence of haemolysis
- High WBC
- Leuco-erythroblastic blood film

Refer to Haematology on urgent (suspected cancer) pathway

If Criteria not met for urgent referral look for causes

If no obvious cause refer to Haematology

Recommended tests pre referral

- FBC
- Viral screen- HIV, EBV
- Autoimmune screen
- Ultrasound abdomen (if not performed)

Other Causes

- Infections – Viral (HIV, EBV, CMV) and parasitic
- Liver: Alcohol, chronic liver disease
- Cardiac failure
- Autoimmune: sarcoid, SLE, rheumatoid arthritis
- Lymphoproliferative disorders
- Myeloproliferative disorders (such as CML or myeloproliferative disorders)
- Haemolysis

Patient Information
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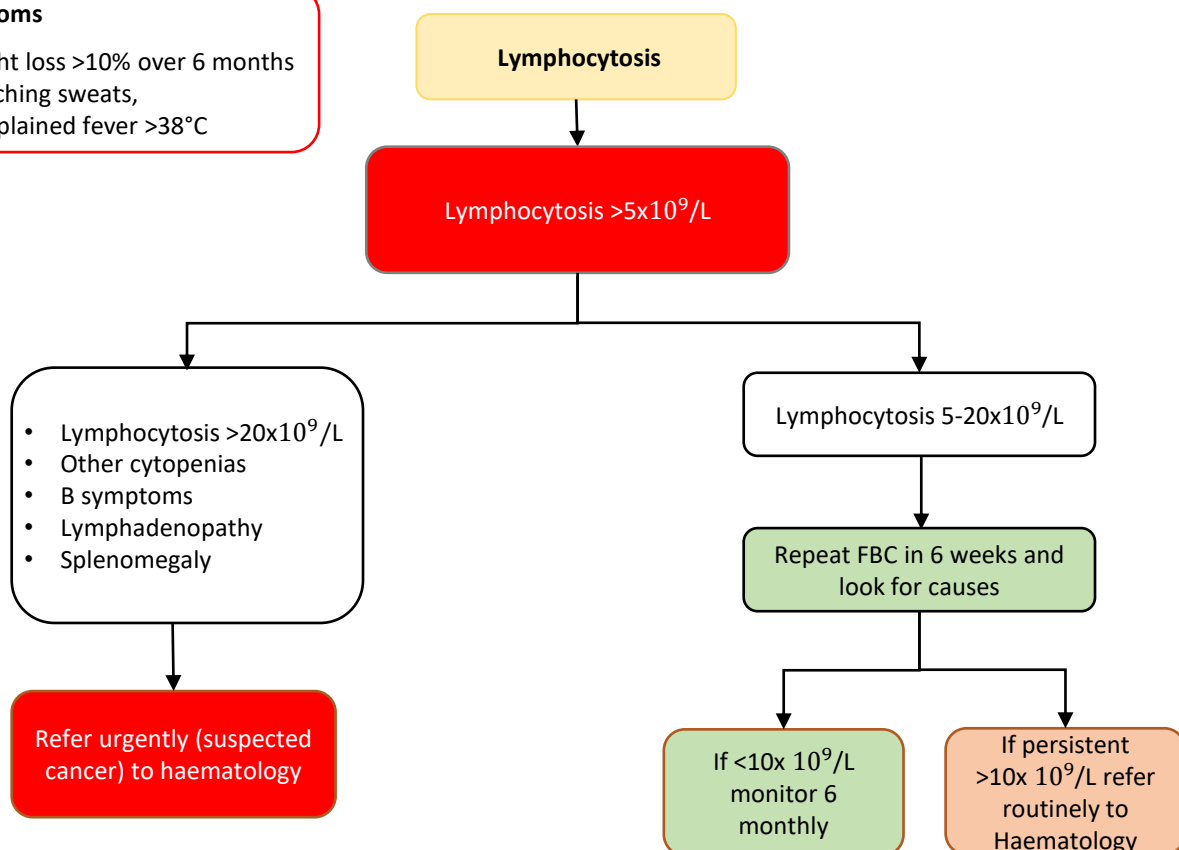
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B Symptoms

- Weight loss >10% over 6 months
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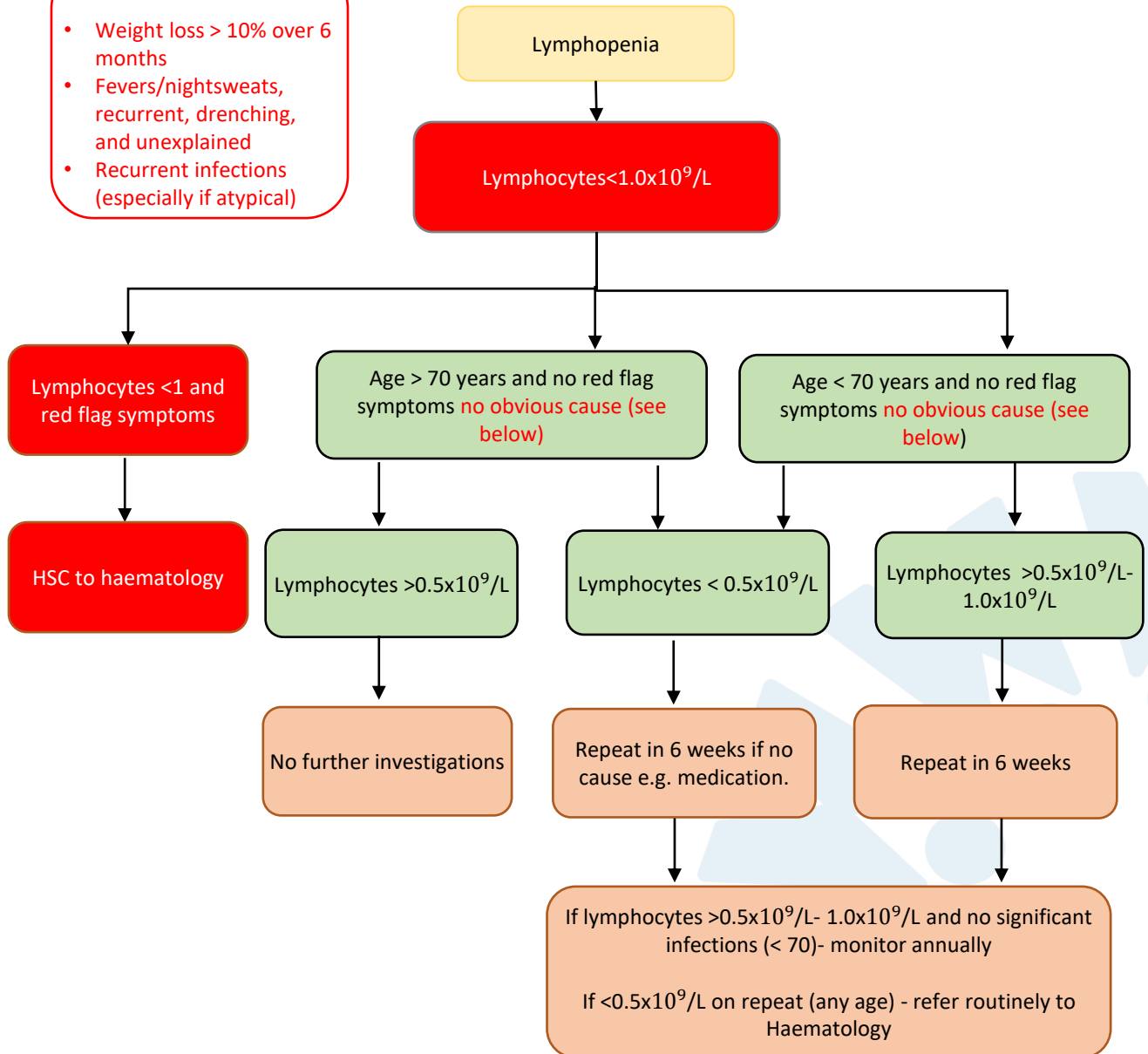


Causes:

- Smoking
- Viral infections especially Glandular fever (EBV)
- Lymphoproliferative disorders (such as CLL)
- Bacterial infections
- Post-splenectomy
- Rheumatoid arthritis

Red Flag signs

- Weight loss > 10% over 6 months
- Fevers/night sweats, recurrent, drenching, and unexplained
- Recurrent infections (especially if atypical)



Causes:

- Elderly patients (>70 years)
- Infections including HIV, hepatitis B and C
- Excess alcohol
- Malnutrition
- Medications: steroids, chemotherapy
- Systemic immune conditions: SLE, rheumatoid arthritis, Sjogren's
- Systemic illness: renal, cardiac, liver failure, malignancy
- Lymphoma

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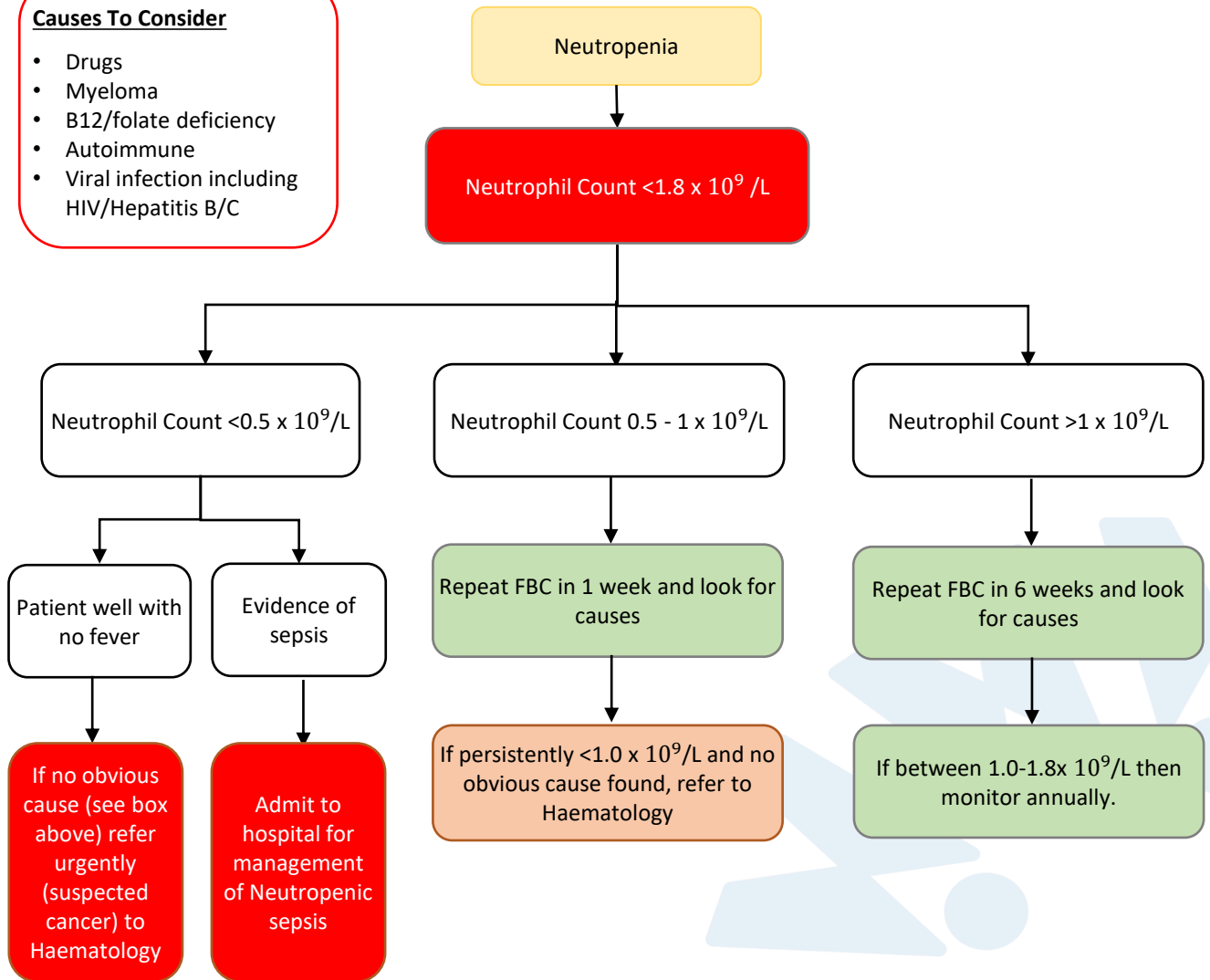
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Causes To Consider

- Drugs
- Myeloma
- B12/folate deficiency
- Autoimmune
- Viral infection including HIV/Hepatitis B/C



Note:

- A neutrophil count of between 1.5 - 2.0 x 10⁹/l whilst below the normal range is unlikely to be of any clinical significance.
- People of Afro-Caribbean or Middle Eastern ethnicity have a lower normal range for the neutrophil count (Duffy Null neutropenia- previously known as constitutional or ethnic neutropenia) 1 - 1.8 x 10⁹/l. This is of no clinical consequence. Only refer if their neutrophils are persistently below **1 for more than 3 months or infective issues.**

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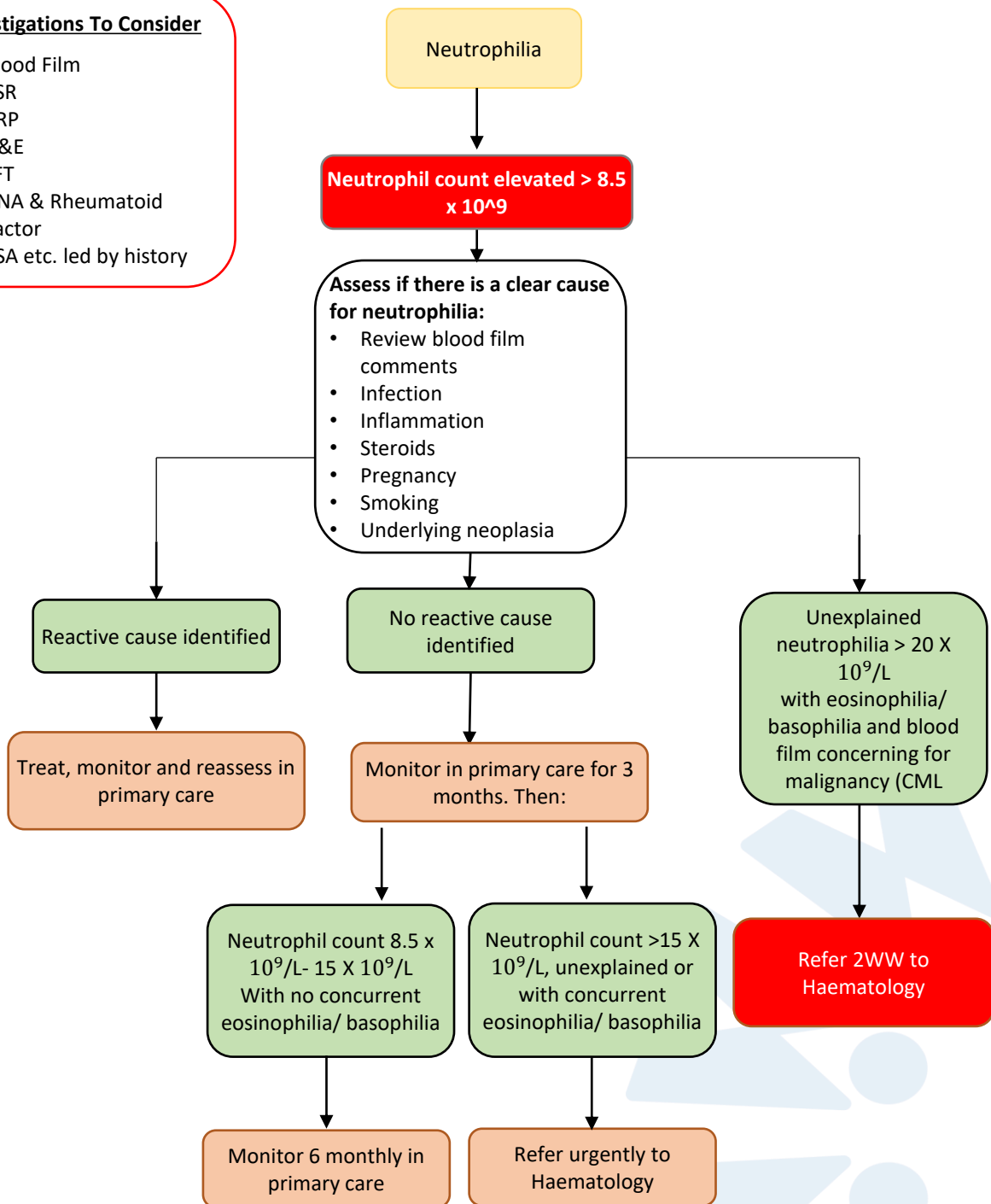
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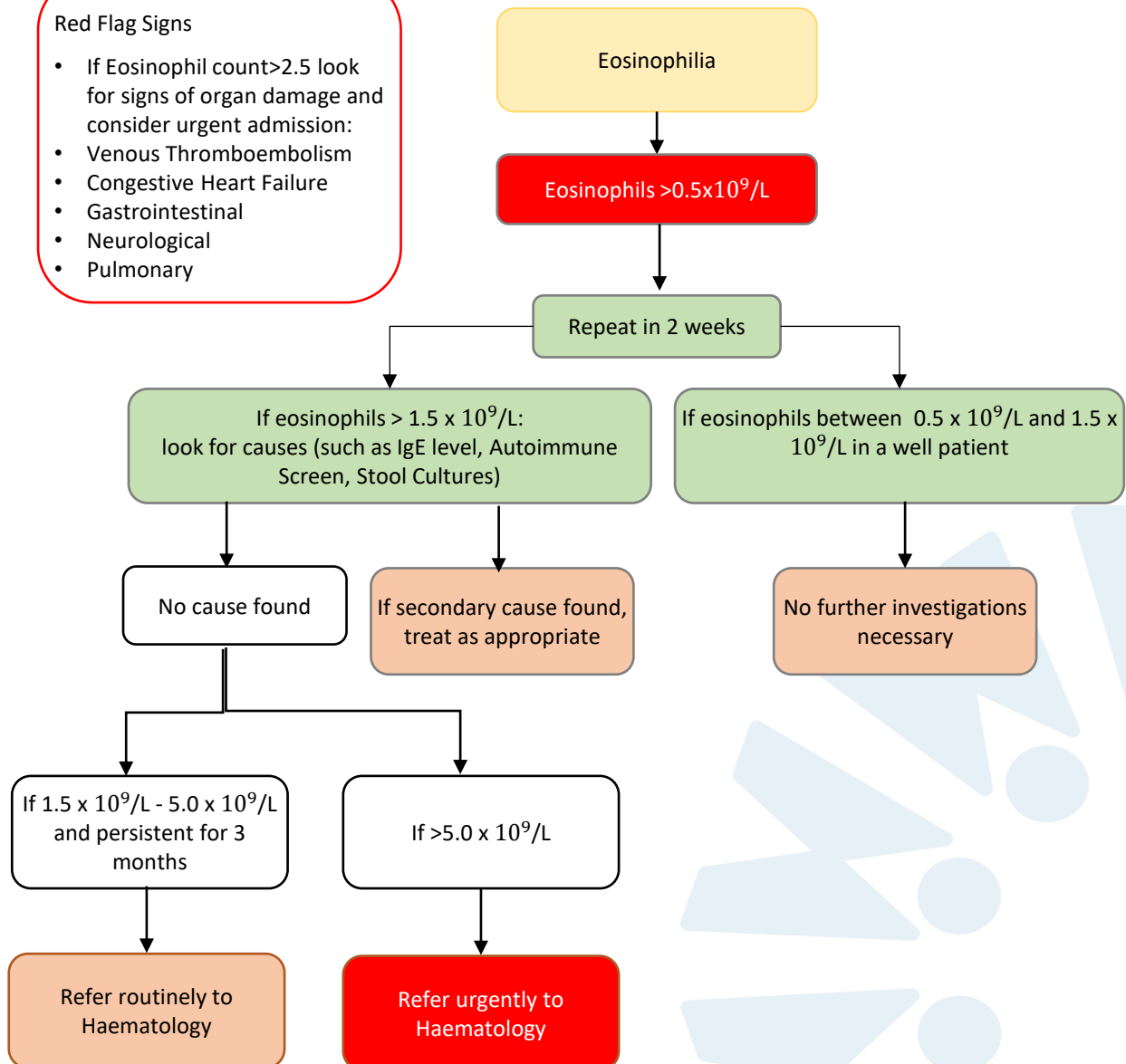
Investigations To Consider

- Blood Film
- ESR
- CRP
- U&E
- LFT
- ANA & Rheumatoid Factor
- PSA etc. led by history



Red Flag Signs

- If Eosinophil count >2.5 look for signs of organ damage and consider urgent admission:
- Venous Thromboembolism
- Congestive Heart Failure
- Gastrointestinal
- Neurological
- Pulmonary



Causes

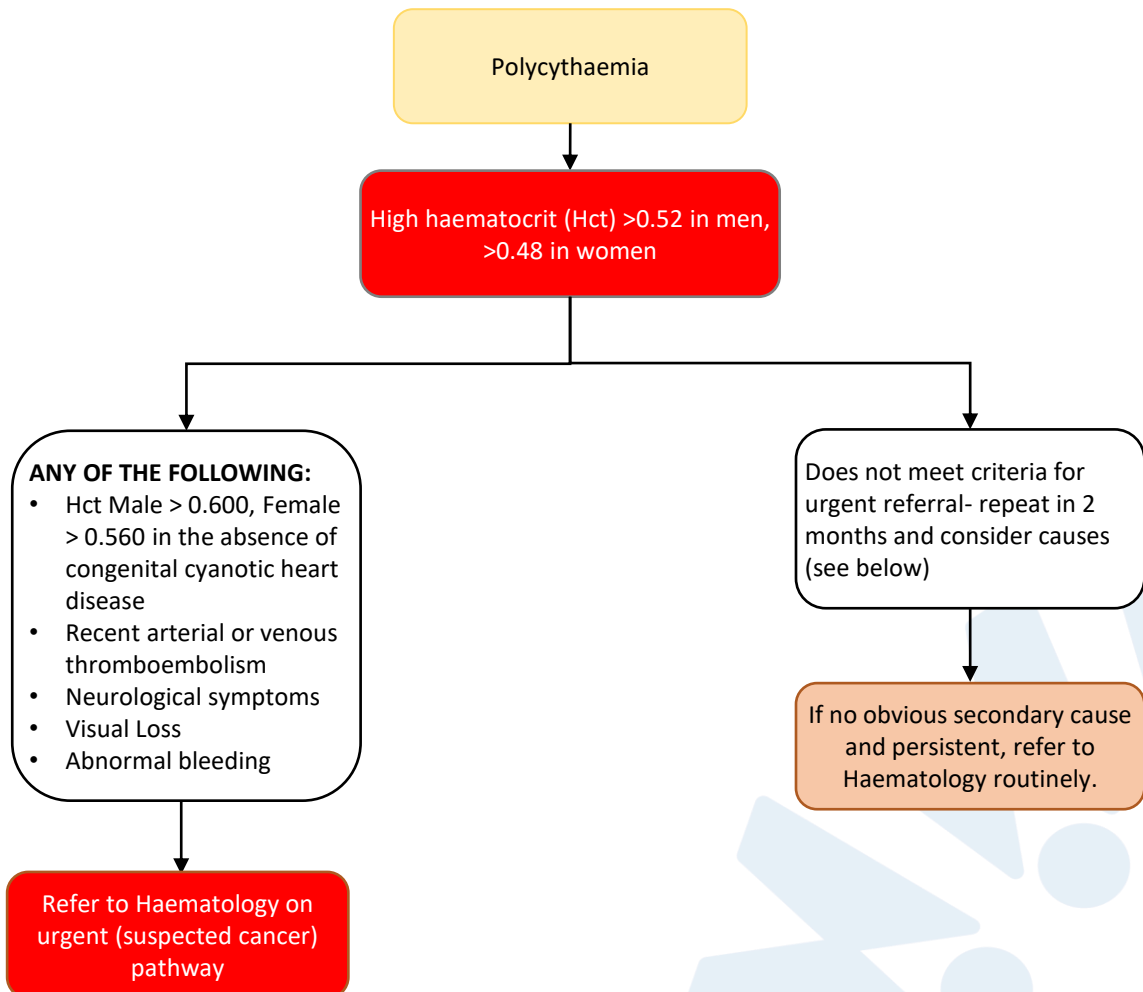
- **Allergy/ atopy:** Asthma / atopic dermatitis / acute urticarial
- **Infections:** especially those due to parasites (most commonly helminthes - hookworm, schistosomiasis - but also giardiasis or other protozoal infections and strongyloides)
- **Drugs:** (penicillins, carbamazepine, sulphonamides are common but any drug is a possible cause)
- **Connective tissue disease** (rheumatoid arthritis, polyarteritis nodosa, Wegener's granulomatosis)
- **Solid malignancy** (breast, renal and lung cancer)
- **Respiratory disease** (Churg-Strauss syndrome, bronchiectasis, cystic fibrosis)
- **Haematological malignancy:** Myeloproliferative disorders

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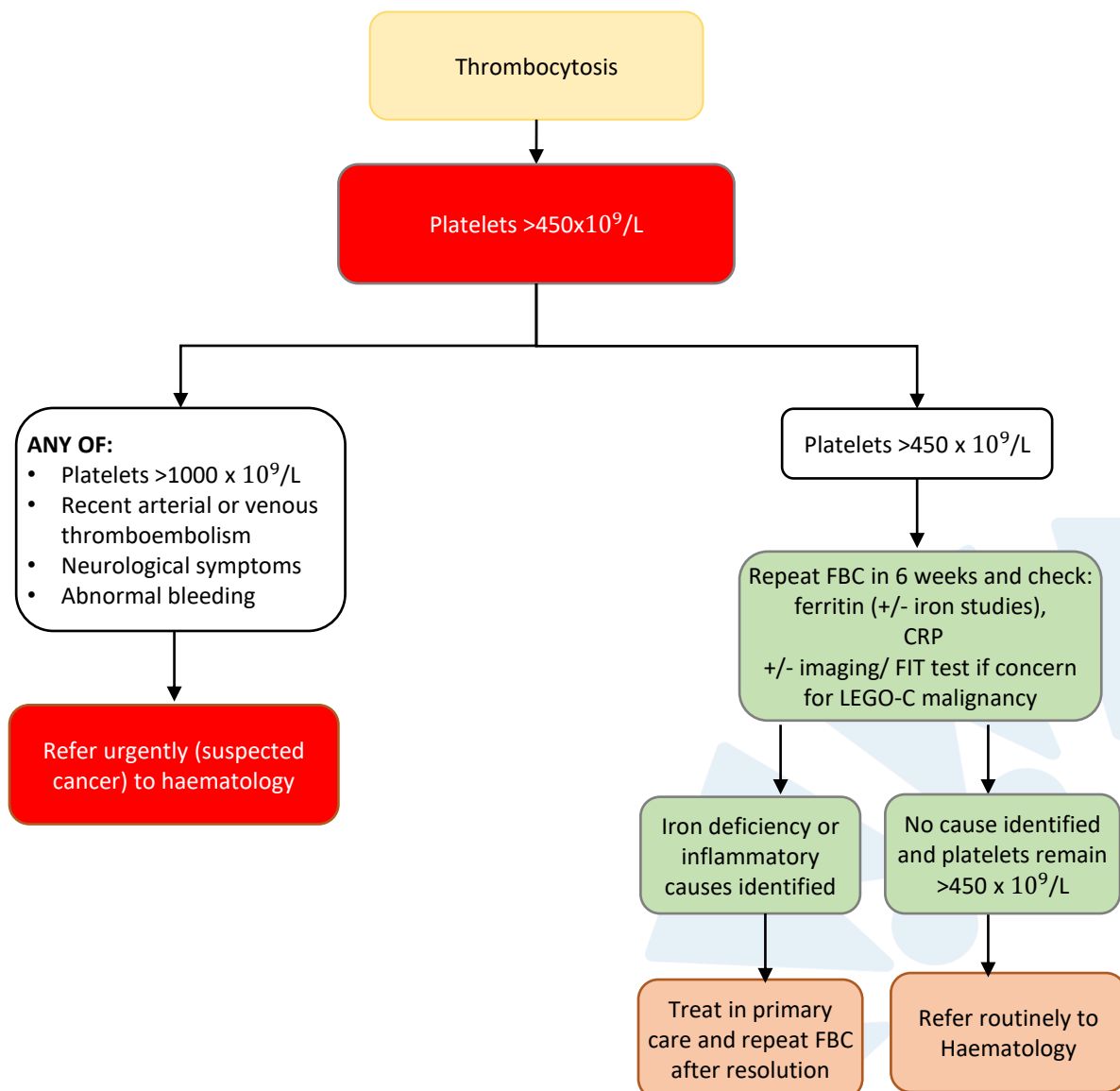
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Causes

- **Drugs** – diuretics, testosterone, anabolic steroids, SGLT2i
- **Lifestyle choices** -smoking, alcohol
- **Hypoxia**- if oxygen sats < 94% in patient with known lung disease (e.g. COPD) consider monitoring in primary care and optimizing medications



Causes

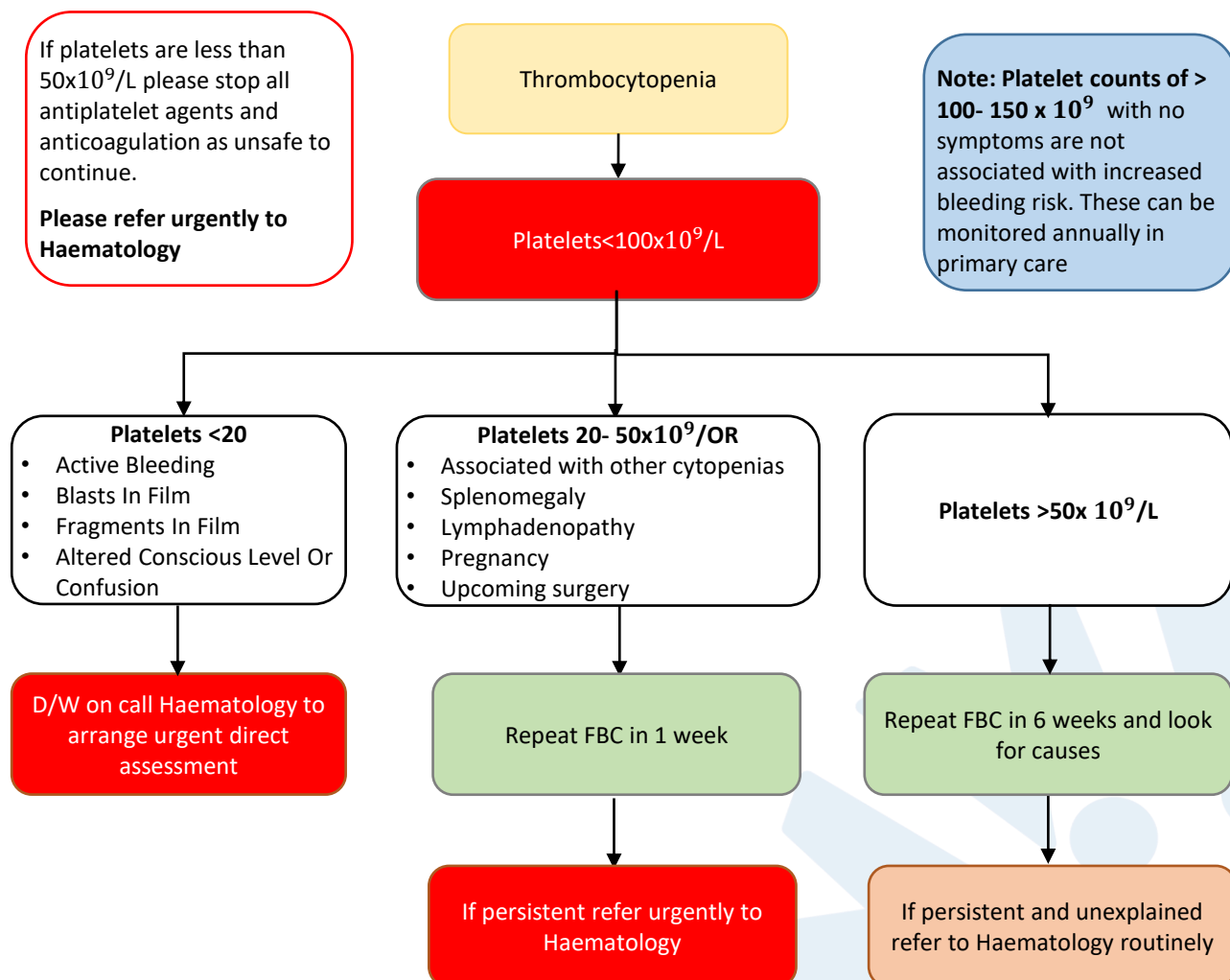
- Iron Deficiency Anaemia
- Inflammation
- Infection
- Post-Splenectomy and Hyposplenism (e.g. Coeliac Disease)
- Myeloproliferative Disorders
- Post-Operatively
- LEGO-C cancers: lung, endometrial, gastric, oesophageal, colorectal

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Causes

- Spurious result from clumping – please look at blood film report and repeat using citrated sample (usually a blue top)
- Immune thrombocytopenic purpura (ITP)
- Alcohol
- Liver dysfunction
- Medications
- B12/folate deficiency
- HIV/Hepatitis B/C
- Bone marrow failure/infiltration

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End Organ Damage (Unexplained):

- Hypercalcaemia (Adj. Calcium > 2.75mmol/L)
- Renal impairment (Creatinine > 177 micromol/L)
- Anaemia (Hb < 100g/L)
- Bone pain (>4-6 weeks), particularly in back/ ribs
- Pathological fracture or lytic lesions on imaging

Paraprotein and/ or abnormal SFLC **RATIO** (see below for normal ranges in renal impairment)

eGFR 45-59: Normal SFLC ratio 0.46-2.62
eGFR 30-44: Normal SFLC ratio 0.48-3.38
eGFR <30: Normal SFLC ratio 0.54-3.30

- IgG paraprotein < 15g/L
- IgM or IgA paraprotein < 10g/L
- Minor SFLC ratio (>0.1 or < 7)

Repeat bloods in 2-3 months

Persistent and stable result

Non-urgent referral to Haematology

Significant rise in paraprotein (>5g/l)

Any of the following:

- IgG Paraprotein >15g/L
- IgM or IgA paraprotein > 10g/L
- IgD or IgE paraprotein
- Significantly abnormal SFLC ratio (<0.1 or > 7)
- End organ damage

Urgent (suspected cancer) referral to Haematology

A polyclonal picture on serum electrophoresis or raised kappa and lambda light chains but with a normal SFLC ratio are reactive features and are not associated with myeloma or MGUS (referral to haematology not indicated)

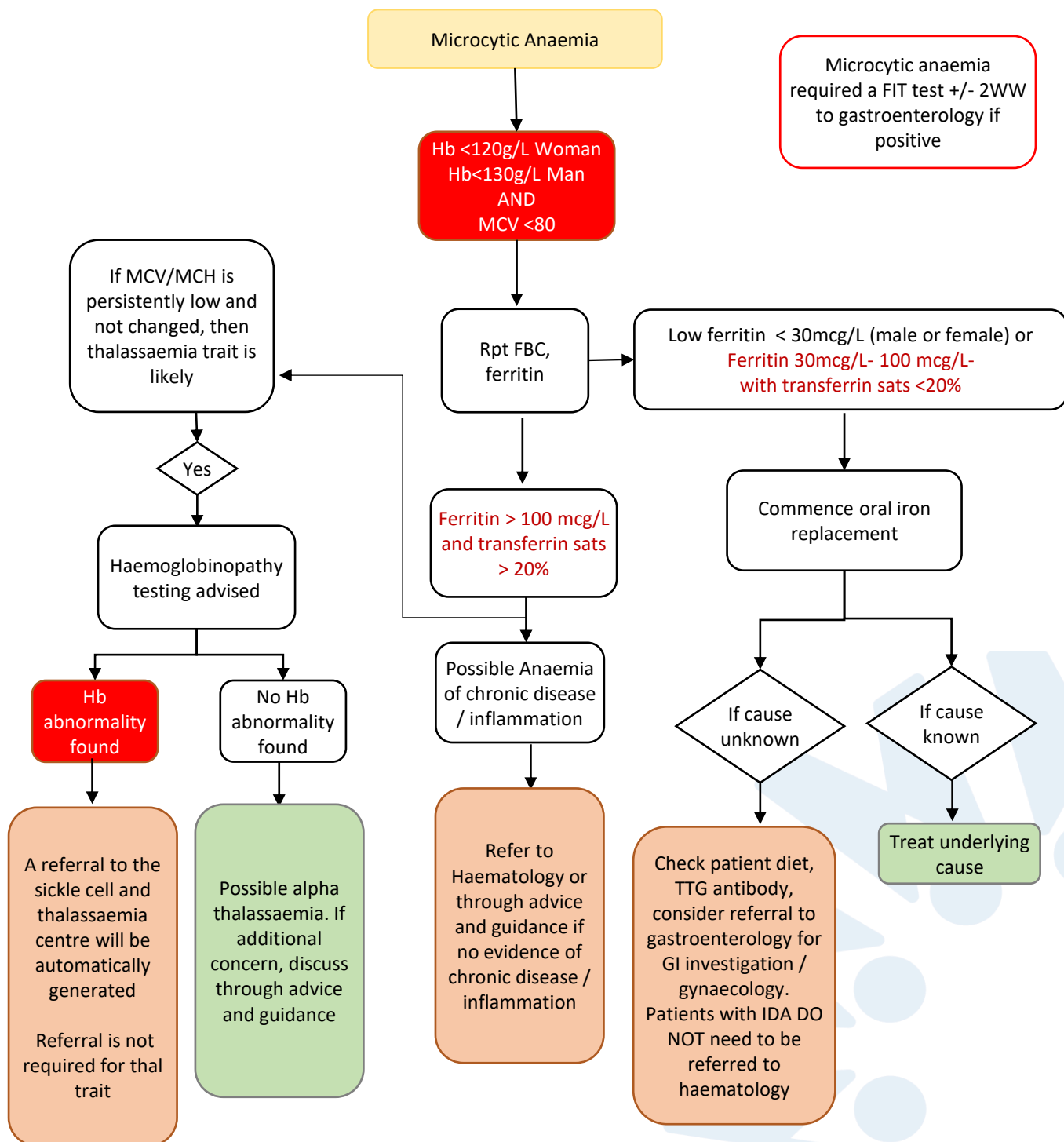
If there are concerns regarding the interpretation of Paraprotein or Serum Free Light Chain results, please discuss with the Haematology team e.g. advice and guidance request

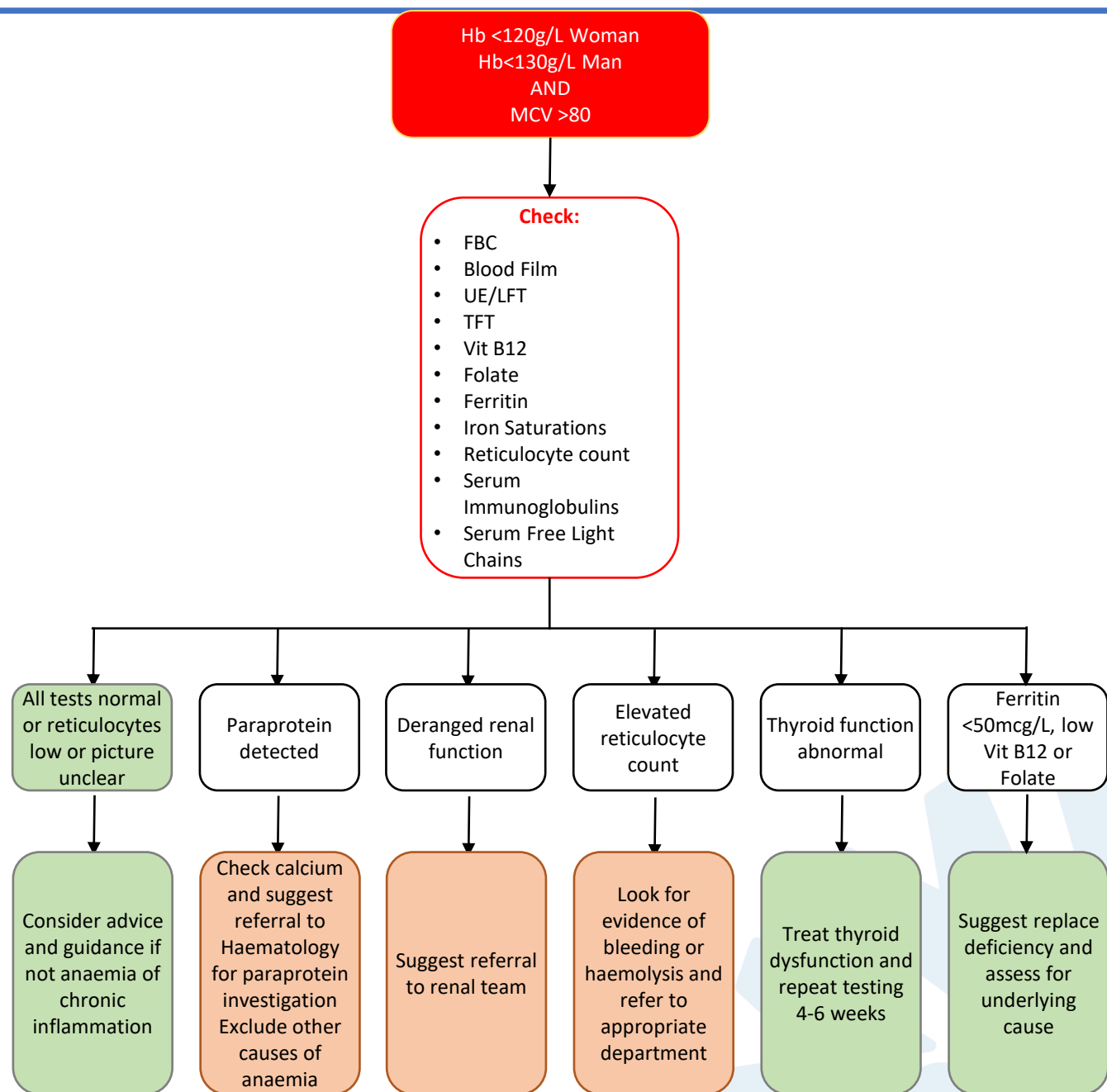
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Note: Markers of haemolysis include a raised reticulocyte count and high bilirubin and LDH.

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Check

- Alcohol history
- Medication (e.g. methotrexate, metformin, some anticonvulsants, hydroxycarbamide, antiretroviral drugs etc.)
- Blood Film
- Vit B12 and folate
- Reticulocyte count/LDH (note often mildly raised, even in absence of clinically significant haemolysis)
- LFT
- TFT
- Serum immunoglobulins
- Serum Free Light Chains
- Family history

Macrocytosis
With/without
anaemia

High Mean Cell Volume
(MCV)*

Repeat FBC to ensure not
spurious (e.g. delayed
transport/ overheating
etc.)

MCV remains
raised

Notes

*A high MCV can be a normal physiological finding in pregnancy

If Vit B12/folate
deficient and Hb
<80g/L or other
cytopenias

Repeat FBC with reticulocyte
count 5-7 days after starting
replacement therapy

Consider referral to Haematology if:

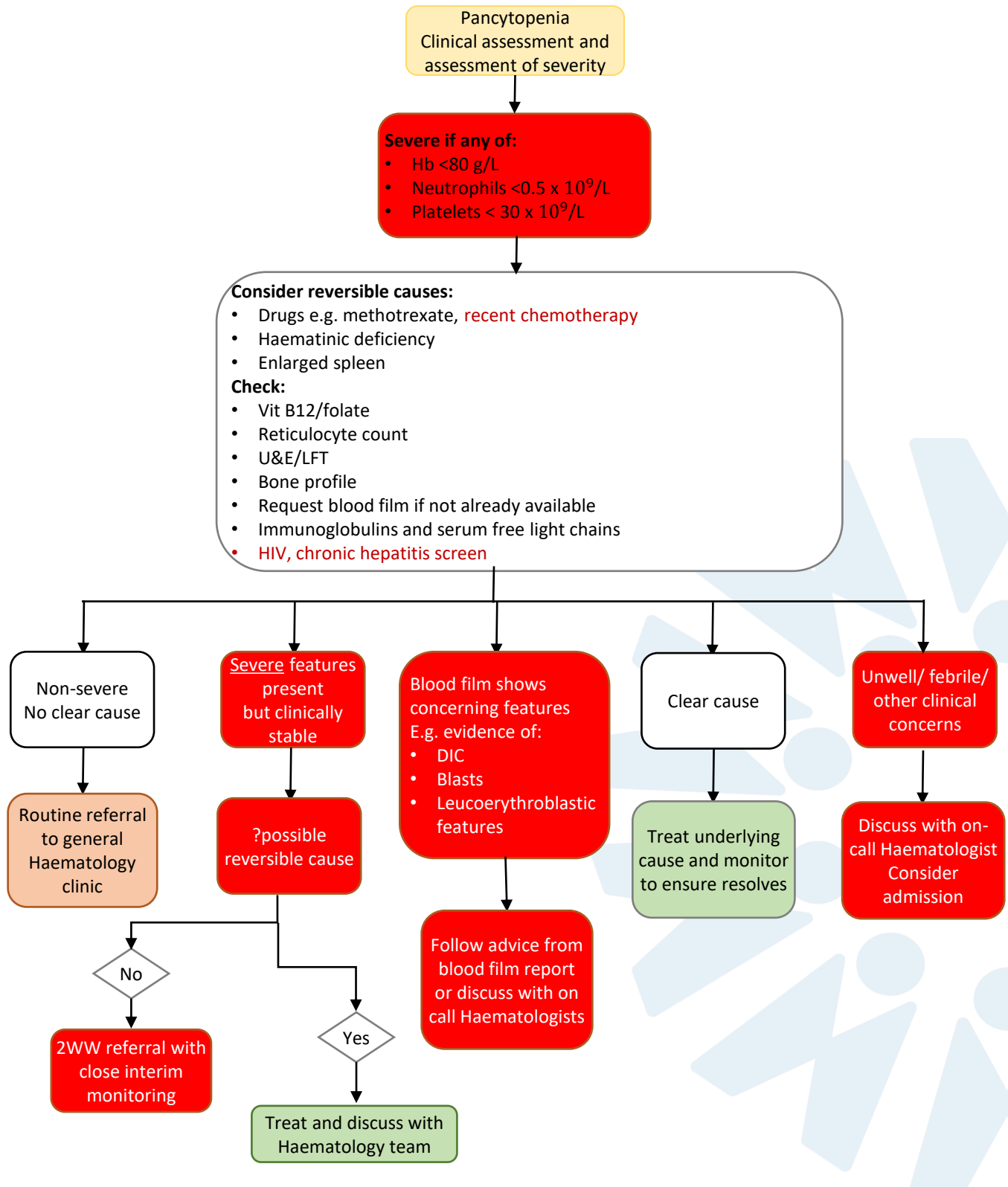
- No secondary cause and MCV >105fL if other cytopenias or >110fL in the absence of other cytopenias
- No history of liver disease
- Dysplasia on blood film
- Paraprotein detected

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***Note: it is recommended to screen ADULT first degree relatives (siblings) of known C282Y HOMOZYGOTES ONLY for genetic haemochromatosis due to their increased risk for C282Y homozygosity. Screening should be performed by iron studies and ferritin, with genetic testing reserved for those with abnormal results. HFE testing can be performed in primary care and does not require referral to haematology or clinical genetics (see text). HFE testing in children is inappropriate as this is an adult onset condition.**

