

**KING'S HEALTH PARTNERS - GSTT QUICK GUIDE TO HAEMATOLOGY**  
**PLEASE ENSURE ALL RELEVANT RESULTS ARE SENT WITH THE REFERRAL**

**Version control:**

There are two controlled versions of this document, one for GSTT and one for KCH. While the clinical content is the same, contact details/suggested clinics differ between the two versions. Any suggested amendments should be submitted to all the document owners:

| GSTT Document Owners   | KCH Document owner  |
|--|---|
| Dr Rachel Kesse-Adu <a href="mailto:Rachel.Kesse-Adu@gstt.nhs.uk">Rachel.Kesse-Adu@gstt.nhs.uk</a> | Dr Robin Ireland <a href="mailto:r.ireland@nhs.net">r.ireland@nhs.net</a> |
| Dr Gulnaz Shah <a href="mailto:Gulnaz.Shah@gstt.nhs.uk">Gulnaz.Shah@gstt.nhs.uk</a>                |   |

Version Control GSTT Document

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|-----------------------------|--|
| <b>Document name</b>        | KHP-GSTT Quick Guide to Haematology          |
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| <b>Version</b>              | Version 1                                    |
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| <b>Owner</b>                | R Kesse-Adu and G Shah                       |
| <b>Author</b>               | Consultant Haematologist                     |
| <b>Approved by, date</b>    | Gulnaz Shah, March 2017                      |
| <b>Superseded documents</b> | Quick Guide to Haematology                   |
| <b>Related documents</b>    | KHP-GSTT Adult Haematology GP Referral Guide |

Version Control KCH Document\*

| Version   | QPulse Document Number  | Change details | Change Owner  | Active date |
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| Version 1 | PDC184: KHP-KCH Quick Guide: Joint Adult Haematology Referral | FINAL DRAFT    | Robin Ireland | May 2018    |

\*Version controlled via KCH Haematology QPulse system: please ensure subsequent revisions of this document are sent to the Haematology Clinical Quality Manager ([Helena.munro@nhs.net](mailto:Helena.munro@nhs.net)). Controlled copy available [Y:\Guidelines](#)

| Differences between KCH and GSTT versions:  | Location        |
|---|-----------------|
| Version control details   | Front page      |
| Suggested clinic to refer to and contact details including suspected spinal cord compression pathway      | In each section |
| Generic comments re Two Week Waits, ConsultantsConnect and Thrombosis/Haemostasis included in KCH version | Front page      |

**QUICK GUIDE TO HAEMATOLOGY PLEASE ENSURE ALL RELEVANT RESULTS ARE SENT WITH THE REFERRAL**

| Referral category  | Suggested tests   | Criteria for urgent referral  | Criteria for routine referral  | Suggested clinic to refer to  |
|--|---|---|--|---|
| <b>Anaemia</b><br>If Iron deficient refer to gastroenterology, gynaecology or urology as appropriate | Detailed history including dietary, blood film, reticulocytes, haematinic assays, immunoglobulins and serum protein electrophoresis to assess for paraprotein, renal function   | Leucoerythroblastic film, unexplained progressive symptomatic, enlarged spleen or lymph glands                        | Persistent unexplained anaemia, intolerance or suboptimal response to oral iron. B12 deficiency + no evidence of pernicious anaemia            | Non-malignant haematology clinic-Guy's site-Haematology-Guy's & St Thomas'                          |
| <b>Erythrocytosis/polythaemia</b><br>Is judged on basis of HCT or PCV.                               | Repeat blood test when non-fasted, alcohol/ smoking, glucose, drugs, JAK2 mutation (97% positive in PVera)  | HCT ♂ >0.60 or ♀ >0.56<br>Recent thrombosis, neurological or visual symptoms  | Persistently elevated HCT ♂ >0.51 or ♀ >0.48 (nb lower if associated iron deficiency). Associated itch or ↑WBC / Plts                          | Myeloproliferative, High Hb and High Platelets-Guy's site-Haematology-Guy's & St Thomas'            |
| <b>Haemochromatosis/Elevated ferritin</b>  | Detailed history and examination- transferrin saturation, HFE genotype, exclude neoplasia, inflammatory markers, Virology-hep B, C, HIV, alcohol status, liver profile, Metabolic syndrome – check BP, BMI, Cholesterol, triglycerides glucose/HBA1C, renal failure, thyrotoxicosis | Evidence of cardiac, liver or endocrine damage  | Persistent unexplained raised ferritin, genetic counselling of relatives.  | Non-malignant haematology clinic-Guy's site-Haematology-Guy's & St Thomas'                          |
| <b>Haemoglobinopathy</b><br>Sickle cell disease and thalassaemia                                     | FBC, Hb Electrophoresis (essential), renal and liver function.  | <b>Acute presentation of severe pain, acute chest syndrome, stroke or priapism</b> should be referred directly to A+E | Sickle cell disease (HbSS, HbSC, HbSB thalassaemia, HbSD, HbSE, HbS-OArab)<br>B thalassaemia major<br>B thalassaemia intermedia<br>HbH disease | Adult Sickle Cell and Haemoglobinopathy clinic-Guy's site-Haematology-Guy's & St Thomas'            |
| <b>Lymphadenopathy</b>   | FBC, blood film, glandular fever, HIV test, monitoring  | >1cm for >6 weeks;<br><6 weeks + B* symptoms;<br>enlarging/>1 site, hepatosplenomegaly, abnormal FBC                  | Persistent lymphadenopathy not meeting urgent criteria   | Suspected Lymphoid Disorders clinic (Lumps & Bumps) - Guy's Site - Haematology - Guy's & St Thomas' |
| <b>Lymphocytosis</b><br>Lymphocytes >4 x10 <sup>9</sup> /L   | Repeat FBC, Blood film, Glandular fever screen if appropriate, smoking history  | Anaemia, ↓ANC, ↓platelets, splenomegaly, painful /progressive lymphadenopathy, B* symptoms                            | Persistent lymphocytes > 5 x10 <sup>9</sup> /L, not meeting urgent criteria  | Suspected Lymphoid Disorders clinic (Lumps & Bumps) - Guy's Site - Haematology - Guy's & St Thomas' |
| <b>Macrocytosis</b><br>Treat B12/folate deficiency before referral. Uncomplicated pernicious         | Blood film B12/folate (IF/coeliac antibodies if abnormal), alcohol/ liver/thyroid screen, serum protein electrophoresis review medications  | Associated neurological symptoms  | Persistent unexplained isolated MCV>105fl or MCV> 100fl and a cytopenia (hb< 100, ↓WBC or platelets < 100).                                    | Non-malignant haematology clinic-Guy's site-Haematology-Guy's & St Thomas'                          |

| Referral category   | Suggested tests   | Criteria for urgent referral   | Criteria for routine referral   | Suggested clinic to refer to   |
|---|---|--|---|--|
| anaemia does not need review  |   |  | Suspected myelodysplasia  |  |
| <b>Neutropenia</b><br><1.5 x10 <sup>9</sup> /L<br>(<0.8 in African Caribbean)   | Review ethnicity + drugs, blood film, haematinics, Virology-hepatitis B and C, HIV, autoimmune screen   | Susceptibility to infection, associated pancytopenia   | Unexplained and persistently <1.5 x10 <sup>9</sup> /L (nb <0.8 in African Caribbean)  | Non-malignant haematology clinic-Guy's site-Haematology-Guy's & St Thomas'                               |
| <b>Eosinophilia</b><br>>1.5 x10 <sup>9</sup> /L   | Blood film<br>Inflammatory markers-CRP and ESR, renal, liver and bone profile<br>LDH, Vitamin B12, Allergy/atopy status<br>Stool cultures for parasites | Leucoerythroblastic film, ANC > 50 x10 <sup>9</sup> /L, AEC > 10 x10 <sup>9</sup> /L, Eosinophils >1.5 x10 <sup>9</sup> /L with evidence of organ damage                         | Eosinophils >1.5 x10 <sup>9</sup> /L  | CML/Eosinophilia Clinic-Guy's site-Haematology-Guy's & St Thomas'  |
| <b>Neutrophilia/leucocytosis</b><br>>15 x10 <sup>9</sup> /L   | Blood film, inflammatory markers, smoking   | Leucoerythroblastic film, ANC > 50 x10 <sup>9</sup> /L,<br><b>Please phone on call haem consultant via switchboard if ANC &gt;100 or Symptomatic for ↑viscosity</b>              | Persistently unexplained WBC >20 x10 <sup>9</sup> /L, Neuts >15 x10 <sup>9</sup> /L   | Suspected Myeloproliferative, High Hb and High Platelets-Guy's site-Haematology                          |
| <b>Paraprotein disorders</b><br>ie presence of monoclonal protein band on serum electrophoresis and/or raised serum free light chains with abnormal ratio and/or presence of urinary Bence jones proteins | FBC, renal and bone profile   | Presence of ↑calcium, ↑lymphs unexplained renal failure, bone pain or pathological #, anaemia, enlarged spleen/lymph glands<br><b>Suspected spinal cord compression by phone</b> | Newly diagnosed paraprotein not meeting criteria for urgent referral                  | Myeloma, amyloid and other paraprotein related disorders - Guy's Site - Haematology - Guy's & St Thomas' |
| <b>Thrombocytthaemia/Thrombocytosis</b><br>Plts >450 x10 <sup>9</sup> /L  | Blood film, exclude iron deficiency with ferritin/iron studies, inflammatory markers  | Plts >1000 x10 <sup>9</sup> /L or >600 recent thrombosis/bleed   | Persistent unexplained plts >450 x10 <sup>9</sup> /L                                  | Suspected Myeloproliferative, High Hb and High Platelets-Guy's site-Haematology-Guy's&St Thomas'         |
| <b>Thrombocytopenia</b><br>Plts <150 (80 in African Caribbean)  | Blood film, repeat for persistence, autoimmune profile, haematinics, liver profile, alcohol history, drug review, HIV, hepatitis B and C test           | Plts <50 x10 <sup>9</sup> /L or 50-100 + other cytopenia, spleen/lymph glands, pregnancy, surgery<br><b>&lt;20 /active bleeding by phone</b>                                     | Persistent <100 x10 <sup>9</sup> /L (<80 in African Caribbean); history of thrombosis | Non-malignant haematology clinic-Guy's site-Haematology-Guy's & St Thomas'                               |

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