

DOCUMENT CONTROL PAGE

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1. Introduction

Children and Young People under the care of the Haematology-Oncology team are at risk of sepsis either because of a central line or neutropenia. This document should be used to risk-stratify children with febrile non- neutropenia into low-risk children who may not need admission or prolonged antibiotics and higher risk children with febrile non neutropenia who need admission and a minimum of 48 hours intravenous antibiotics.

This document should be used in conjunction with "Paediatric Haematology/Oncology Antibiotic and Antifungal Treatment Guidelines".

These guidelines do not apply to patients receiving a stem cell transplant or those with a benign haematology condition

2. Purpose

This guidance is produced in response to NICE CG 151 (Sep 2012). It should be read in conjunction with "Paediatric Haematology/Oncology Antibiotic and Antifungal Treatment Guidelines"

http://microbiology.staffnet.xcmmc.nhs.uk/media/400809/paediatric_haematology_oncology_antibiotic_and_antifungal_guidelines_june_2017_pdf.pdf

This guidance is designed primarily for use in the Haematology and Oncology department and PED units at Royal Manchester Children's Hospital, RMCH. The principles of practice are applicable to Paediatric Oncology Shared Care Units (POSCUs).

3. Roles and Responsibilities

- 3.1 The Clinical Team in Malignant Haematology and Oncology are responsible for ensuring that they are familiar with this policy. It is the role of the attending consultant to review admitted patients. Senior staff should ensure this policy is covered as part of induction of new staff members.
- 3.2 The Clinical Team in PED are responsible for ensuring that they are familiar with this policy.

4. Detail of Procedural Document.

This guidance covers:

- 4.1 Definitions of fever, non neutropenia, low risk and higher risk patient populations
- 4.2 Initial management of patients (PED/POSCU/Day Case)
- 4.3 Management of low-risk patient population (low risk disease AND low risk clinical status)
- 4.4 Management of higher risk patient population
- 4.5 Ongoing management statement

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- 4.1 Definitions of fever, low risk and higher risk patient populations
 - 4.1.1 Fever or Febrile is defined as single temperature >38°C
 - 4.1.2 **NON-Neutropenia** is defined as neutrophil count >0.5
 - 4.1.3 **Low risk patients**: as defined by disease AND clinical status. All other patients should be considered **higher risk**.

		<u>Higher risk</u>
<u>Underlying</u> <u>disease</u>	 Acute Lymphoblastic Leukaemia (ALL) Infant ALL on maintenance All oncology patients CNS or non- 	Acute myeloid leukaemia (AML) Infant ALL pre-maintenance
Clinical	CNS •	• Ago <6 months
<u>Clinical</u> <u>status</u>	 Age >6 months No evidence of septic shock No evidence of dehydration Normal neurology 	 Age <6 months Septic shock Vomiting or abdominal pain Reduced urine output or reduced skin turgor or poor feeding in infants > 6 months Altered mental status or focal neurology Requiring opiate analgesia for mucositis Social reasons requiring inpatient management or significant parental concern

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4.1.4 definition of septic shock

Septic shock is defined if the capillary refill >2 seconds, or have been given a fluid bolus, or if systolic BP <5th centile (see chart below, hypotension is a pre-terminal sign in children). If in doubt, discuss with on call Consultant Haematologist/Oncologist.

Age (Years)	Systolic BP (mmHg) 5 th Centile	Systolic BP (mmHg) 50 th Centile
<1	65-75	80-90
1-2	70-75	85-95
2-5	70-80	85-100
5-12	80-90	90-110
>12	90-105	100-120

Ref: Advanced Paediatric Life Support: A practical approach to emergencies. 5th Ed. 2016

4.2 Initial management of patients (PED/POSCU/Day Case)

- 4.2.1 All Haematology and Oncology patients with a single temperature >38°C AND a central line/Portacath in situ should receive a first dose of antibiotics according to the "Paediatric Haematology/Oncology Antibiotic and Antifungal Treatment Guidelines"
- **4.2.2** Patients should then be subsequently assessed by Haematology and Oncology team following the risk stratification outlined in this document and managed following the appropriate pathway. **Febrile and neutropenic patients should be treated as per High Risk patients.**

4.3 Management of low-risk patient population (low risk disease AND low risk clinical status)

- 4.3.1 Line, and urine cultures must be taken.
- 4.3.2 **Low risk patients** who are **non-neutropenic** do not require admission and can be discharged by a member of Haematology or Oncology medical team (or Senior Member of the medical team at POSCU centres). Oral antibiotics may be commenced at discretion of reviewing team if a bacterial focus for infection is suspected.
 - 4.3.2.1 Discharged patients should be added to the Haematology/Oncology handover sheets, maintained by the ward medical team. It is the responsibility of the attending consultants to ensure patients and contact details are added to handover sheets and to follow up on microbiology results. In POSCU centres this role should be allocated to the Senior Member of the medical team.

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4.3.2.2	On discharge parent/carer is advised to come back to hospital (PED) or
	POSCU centre if child becomes more unwell i.e. follow the pathway for
	unwell child relevant for specific centre.

- 4.3.2.3 The ward attending Haematology, Oncology or POSCU medical team must review any cultures at 24 and 48 hours for patients under their care.
- 4.3.2.4 Parent/carer should be contacted by ward team at specific centre at 48 hours with culture results and telephone consult with the attending Haematology, Oncology or POSCU medical team.
- 4.3.2.5 If the patient remains febrile at 48 hours, the patient should come back to hospital (PED) or POSCU centre for further observations, FBC and medical review. Likewise, if blood cultures are positive they need to come back to hospital (PED) or POSCU for review.

4.3 Management of higher risk patient population

- 4.3.3 All patients stratified as higher risk must be admitted for IV antibiotics as per "Paediatric Haematology/Oncology Antibiotic and Antifungal Treatment Guidelines".
- 4.3.4 All higher risk patients need to remain in hospital until cultures are confirmed as negative at 48 hours AND have been afebrile for 24 hours.
- 4.3.5 Ensure review of antibiotics at 48 hours as per "Paediatric Haematology/Oncology Antibiotic and Antifungal Treatment Guidelines".
- 4.3.6 Cultures should be repeated every 48 hours if patient remains febrile.

4.4 Ongoing management statement

- 4.4.1 Patients with ongoing fever post 48 hours should be managed according to the "Paediatric Haematology Oncology Antibiotic and Antifungal Treatment Guidelines"

 http://microbiology.staffnet.xcmmc.nhs.uk/media/400809/paediatric_haematologyoncolog_antibiotic_and_antifungal_guidelines_june_2017_pdf.pdf
- 4.4.2 Patients with positive blood culture must be discussed with the Microbiology Department (RMCH or POSCU where relevant).
- 4.4.3 Patients with suspected viral infections (e.g. VZV, HSV) should be discussed with Virology colleagues at RMCH.

5. Equality Impact Assessment

An initial Equality Impact Assessment has been completed for this document (please refer to the Document Control Page).

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6. Consultation, Approval and Ratification Process

Consultation: This guidance has been drawn up as a joint venture with representation from Haematology, Oncology, PED, POSCU centres and Pharmacy.

Approval and Ratification: RMCH Quality and Safety Committee.

7. Dissemination and Implementation

7.1 Dissemination

This guidance will be publicised and circulated on Ward 86, Ward 84 OPD, PED and POSCU centres. It will be posted on the Intranet website. It will be publicised to the Haematology and Oncology medical and nursing staff, including face to face training at induction.

7.2 Implementation of Procedural Documents
Implementation will be audited 12 months after launch.

8. Monitoring Compliance of 'Guidance for risk stratification of febrile neutropenia and non-neutropenia in Haematology and Oncology'

Compliance will be monitored by cyclical audit at appropriate time frequency.

Process for Monitoring Compliance and Effectiveness

The Lead Consultant for this document is responsible for monitoring compliance with the 'Guidance for risk stratification of febrile neutropenia and non-neutropenia in Haematology and Oncology' at Division and Corporate Level.

This will be completed on a one- to two yearly basis and reported to the Trust Audit Committee.

The following will be monitored for compliance:

Adherence to guidance outlined in document via audit.

Any shortfalls identified will have an action plan put in place to address which will have timescales included for re-audit/monitoring.

9 Standards and Kay Performance Indicators 'KPIs'

All patients for whom this document is relevant should receive treatment in accordance with Section 4 of the guidelines, which will provide a basis for audit as above.

This Policy must be reviewed at least every three years or when there are significant changes to the document or procedure.

Awareness of the policy will be delivered at RMCH Quality and Safety Committee, the local CSU Clinical Effectiveness meeting and the Greater Manchester Cancer Pathway Board meetings for POSCU awareness.

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10 References and Bibliography

Neutropenic sepsis: prevention and management in people with Cancer, National Institute for Clinical Excellence document CG151, September 2012

11 Associated Trust Documents

"Paediatric Haematology/Oncology Antibiotic and Antifungal Treatment Guidelines" http://microbiology.staffnet.xcmmc.nhs.uk/media.400809/paediatric_haematologyoncology_antibiotic_and_antifungal_guidelines_june_2017_pdf.pdf

12 Appendix

Summary Guidance for Risk Stratification of febrile (T>38) Neutropenia (N<0.5) and Non-Neutropenia in Haematology and Oncology.

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SUMMARY GUIDANCE FOR RISK STRATIFICATION OF FEBRILE (T>38) NEUTROPENIA (N<0.5) AND NON-NEUTROPENSIA IN HAEMATOLOGY & ONCOLOGY

