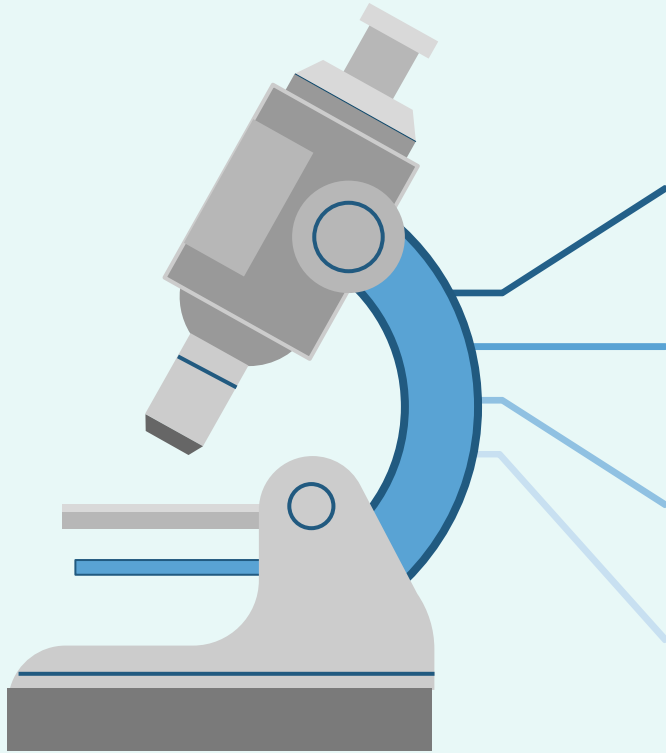




The wonderful World of Blasts!

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Session Aims



1

The anatomy of a blast.

2

When do we see blasts?

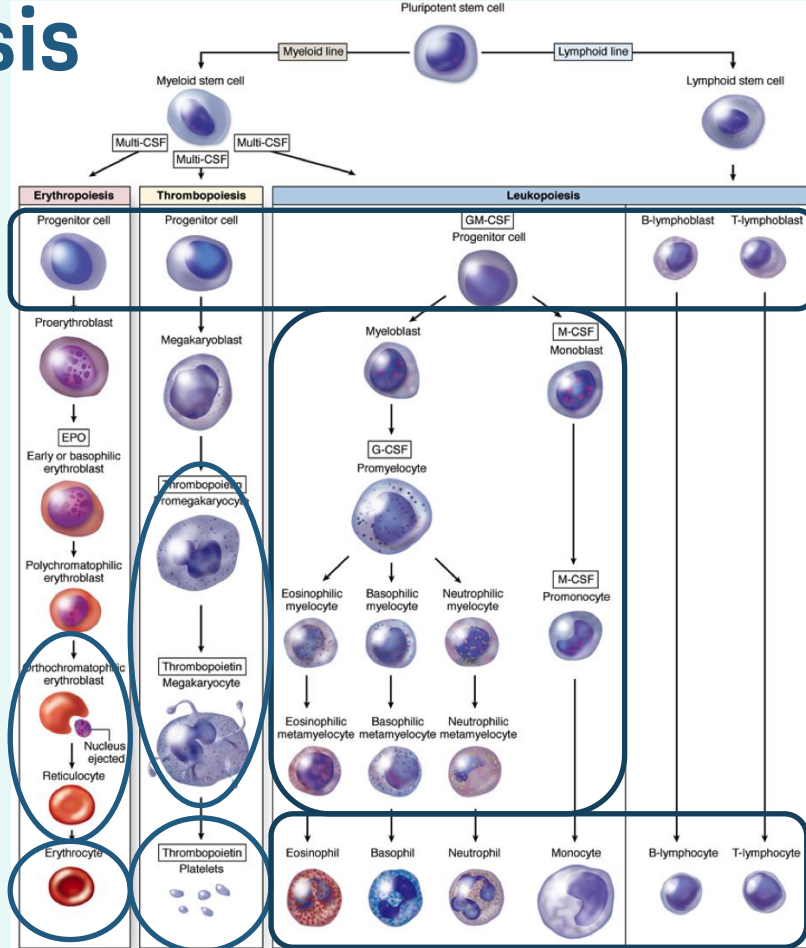
3

When is a blast not a blast?

4

When are blasts important?

Haematopoiesis



Anatomy of a Blast

All blood cells are made in the bone marrow, but only a few enter into peripheral blood (PB).

Megakaryocytes are extremely uncommon in PB.

Red cell precursors are common in anaemia

These cells are normal in peripheral blood.

Blasts are always concerning in peripheral blood!

It can be very hard to tell what kind of blasts they from appearance alone!

Myeloid precursors can be present in PB for many reasons- not all are malignant!

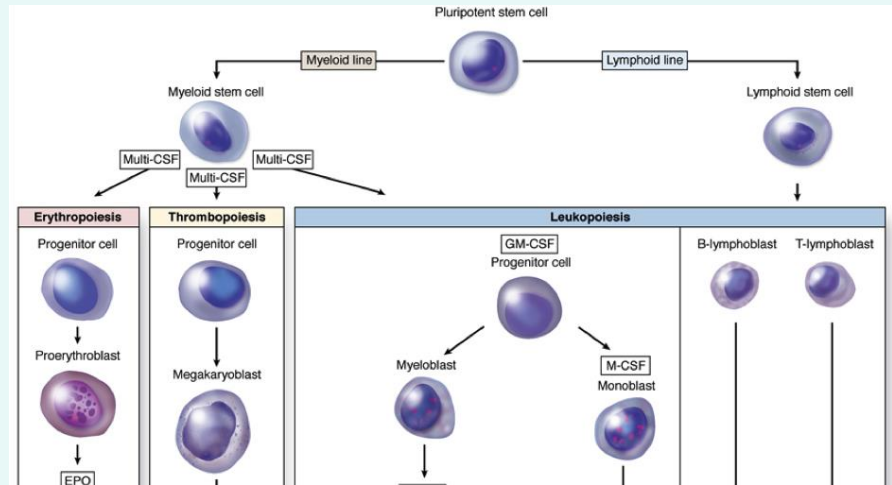
What is a blast?

Blasts encompass a variety of different early stem cells.

Development is driven by key cytokines e.g. GCSF, EPO, IL-6.

As cells develop they gain features which help to identify them.

This may include size, granules, cytoplasm colour, cell markers etc.



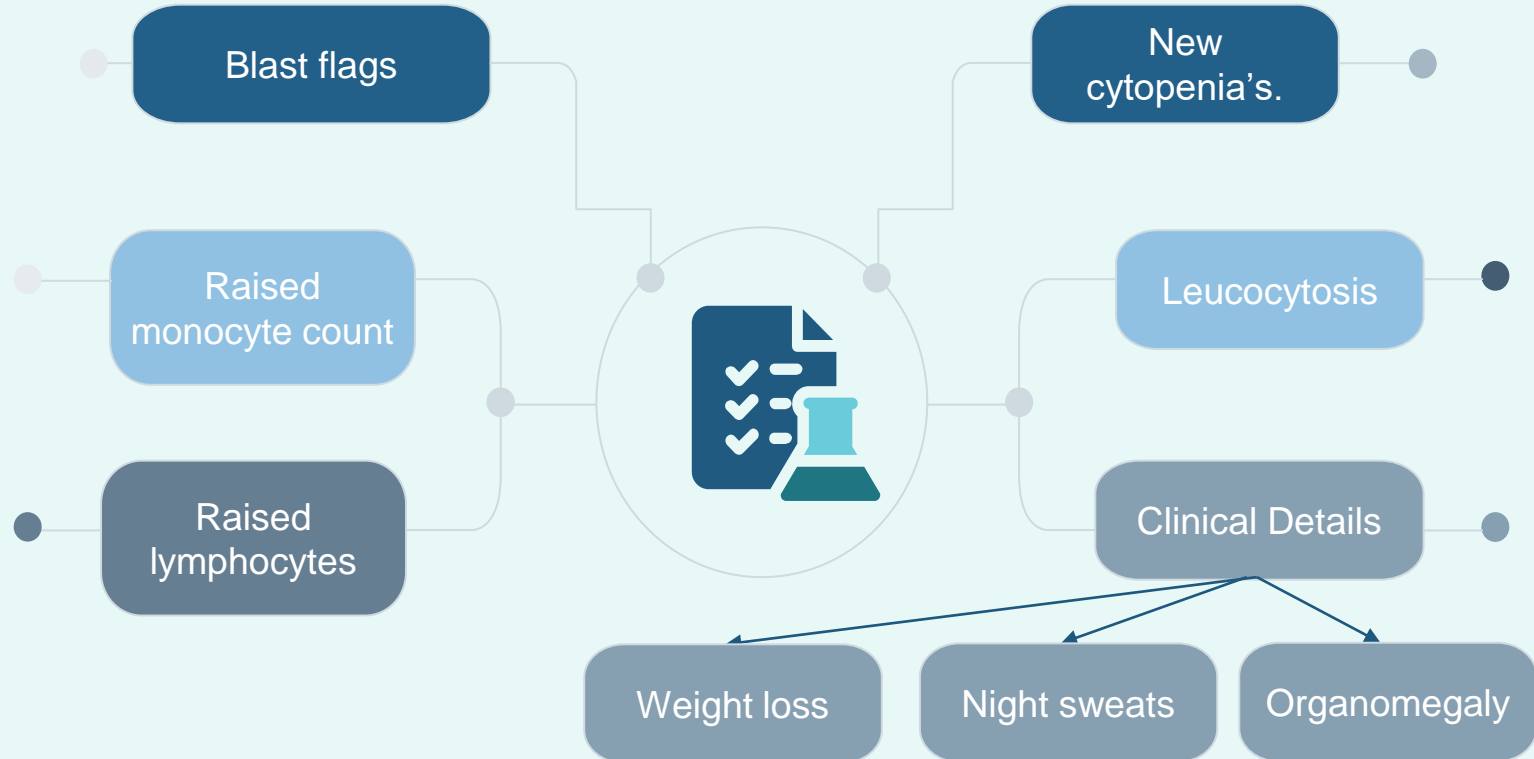
All haematological cells originate from pluripotent stem cells.

Blasts are normal in the marrow in small numbers

Genetic aberrancies can cause maturational arrest, resulting in increased blasts.

What FBC features make us suspicious?

Anatomy of
a Blast.



The Anatomy of a Blast.

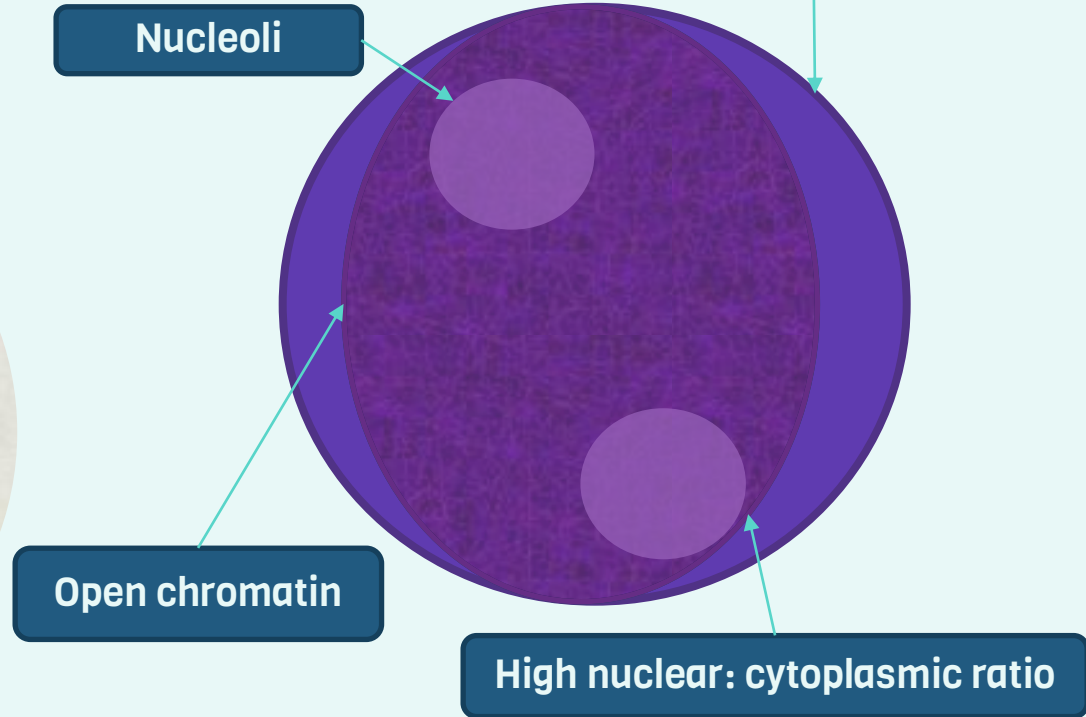
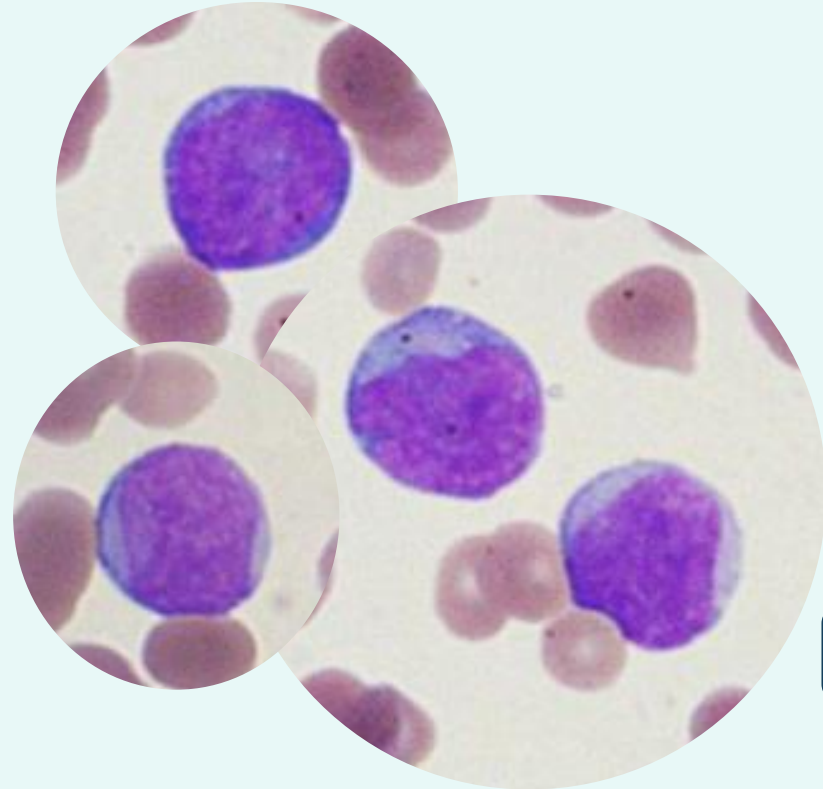
Anatomy of a blast

Basophilic Cytoplasm

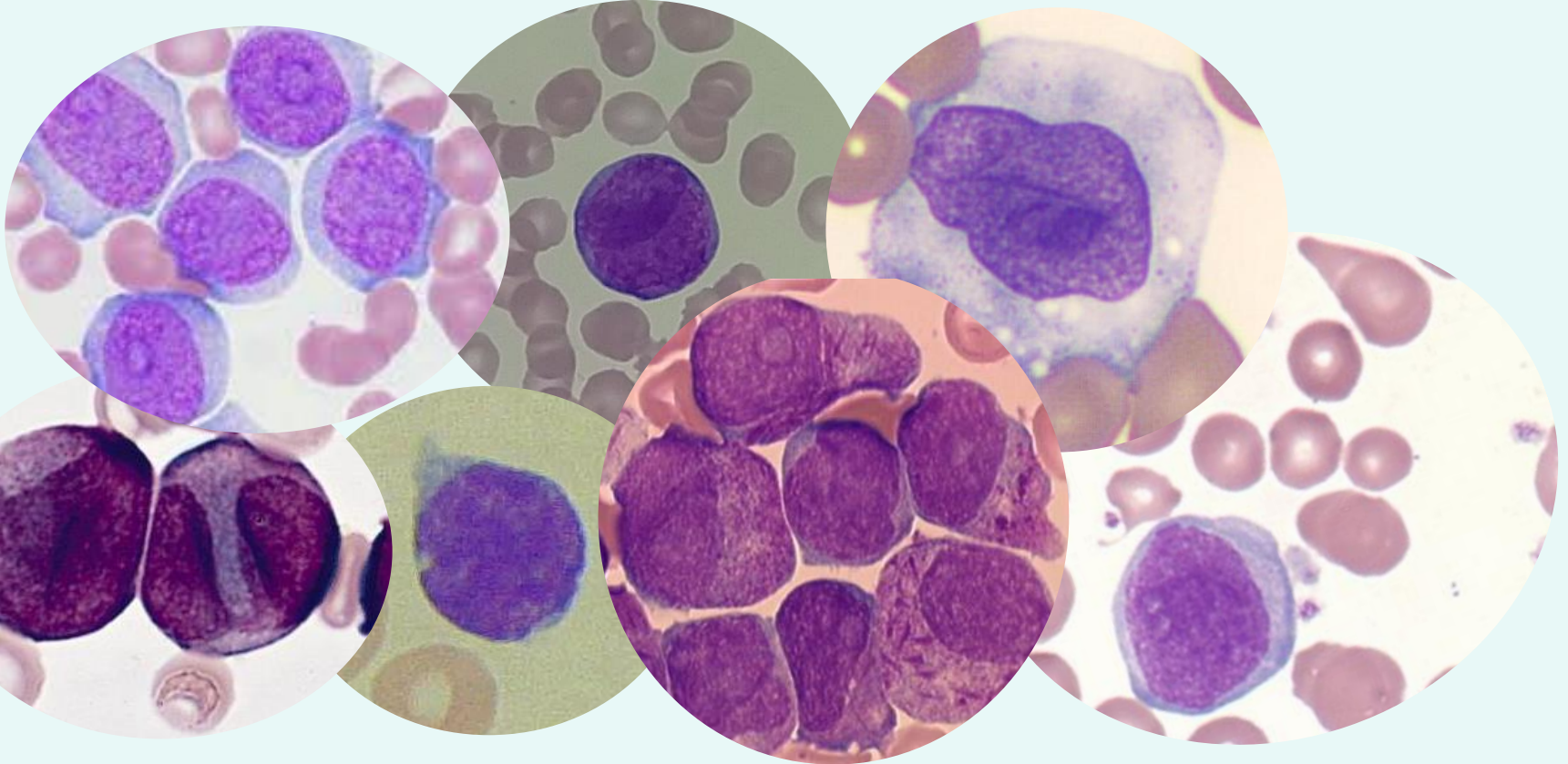
Nucleoli

Open chromatin

High nuclear: cytoplasmic ratio



So all Blasts are the same, right?

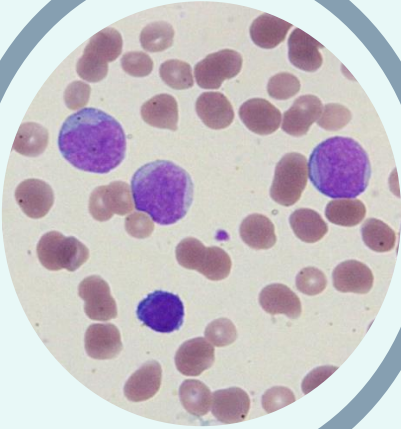




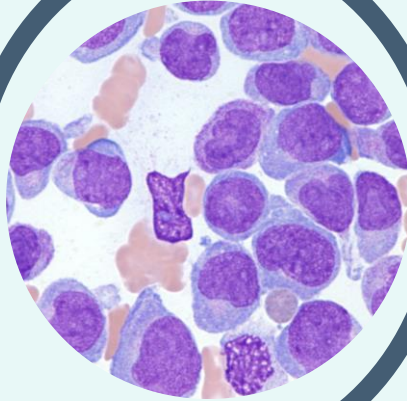
**When do we see
Blasts in Blood?**

Leukemic Causes: Acute

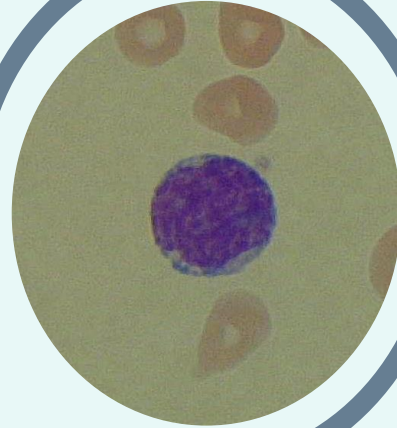
When do we see blasts?



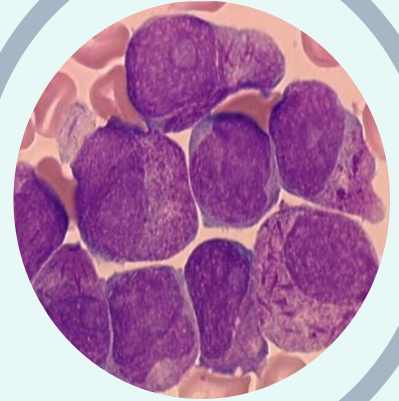
Acute Myeloid
Leukaemia



Acute
Myelomonocytic
Leukaemia



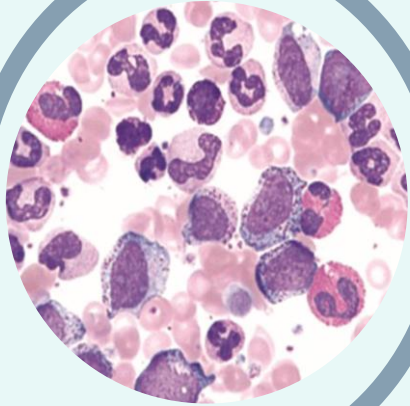
Acute
Lymphoblastic
Leukaemia



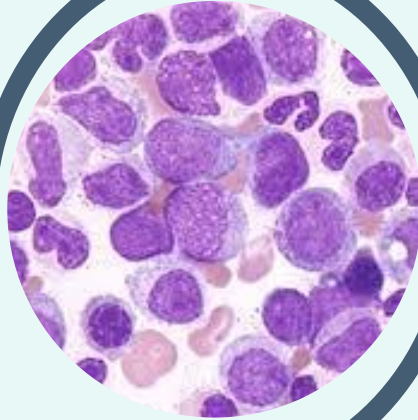
Acute
Promyelocytic
Leukaemia

Leukemic Causes: Chronic

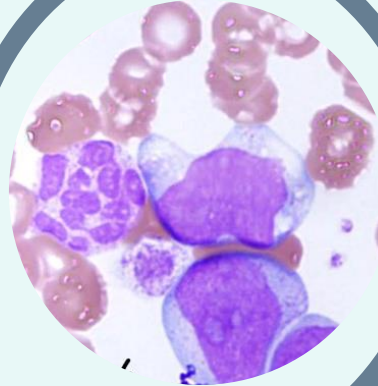
When do we see blasts?



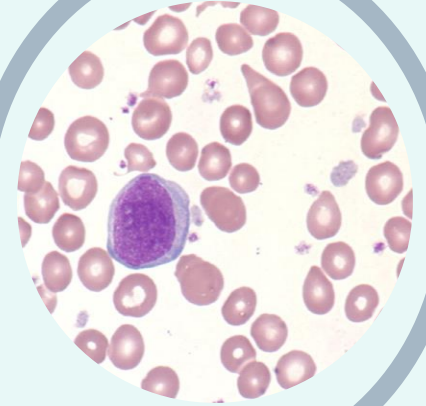
Chronic Myeloid Leukaemia



Chronic Myelomonocytic Leukaemia



Myelodysplastic Syndrome



Myelofibrosis

Myelodysplasia and Blasts

When do we see blasts?

MDS classifications are discrete, but can have a spectrum-like presentation or pattern of development.

MDS-MLD

2-3 lineages affected.

Peripheral blasts: <1%
Marrow blasts: <5%

MDS with excess blasts (MDS-EB) are considered high risk, but patient's may not progress to AML for many years.

MDS-EB2

1+ lineages affected.

Peripheral blasts: 5-19%
Marrow blasts: 10-19%

MDS-SLD

1 lineage affected.

Peripheral blasts: <1%
Marrow blasts: <5%

There are other categories of MDS, but in these cases, blasts are <5% in the bone marrow.

MDS-EB1

1+ lineages affected.

Peripheral blasts: 2-5%
Marrow blasts: 5-10%

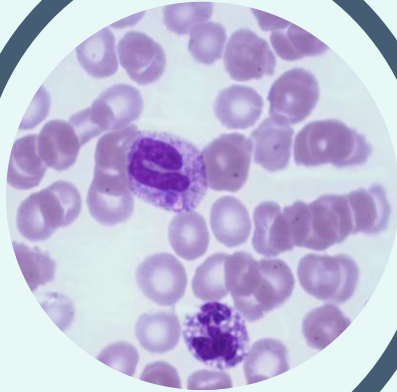
Once blasts are >20% in the bone marrow, MDS becomes AML. PB blasts may increase and cytopenia's may develop/worsen.

Non-Leukemic Causes

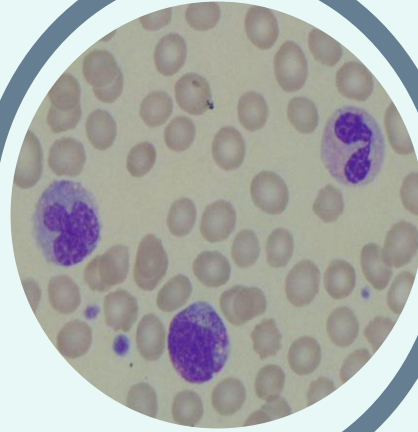
When do we see blasts?



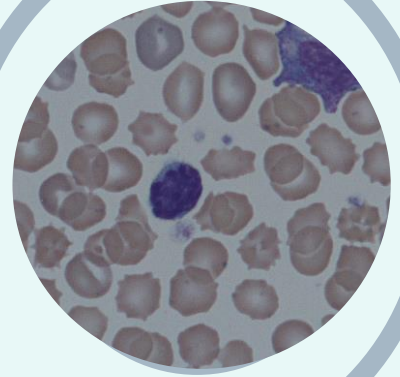
Marrow Infiltration



Bacterial Sepsis



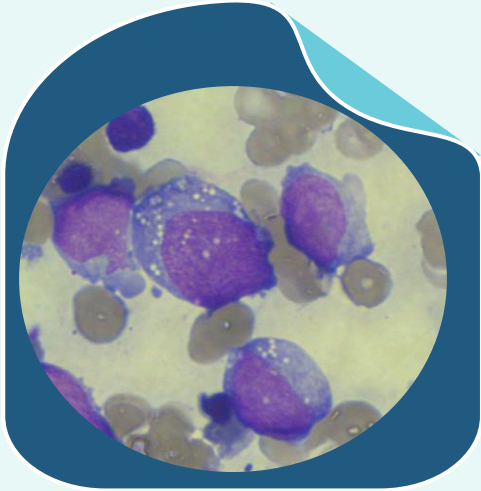
G-CSF



Prematurity

When is a blast
not a blast?

When is a blast, not a blast?



High Grade Lymphoma

Large cells with blastic appearance and vacuolation. These are not blasts- but these patients do need urgent treatment!



EBV/CMV Infection

Cytoplasm can become expansive and dark and nucleoli present, but chromatin will remain clumped!



Prolymphocytes

Prolymphocytes can be seen in LPDs e.g. CLL- but >55% indicates PLL! These cells have nucleoli, but an otherwise mature appearance!

When are blasts important?

When should results be passed to clinicians?

To some extent, all of these are the right thing to do- it all depends on context!

whenever anything changes

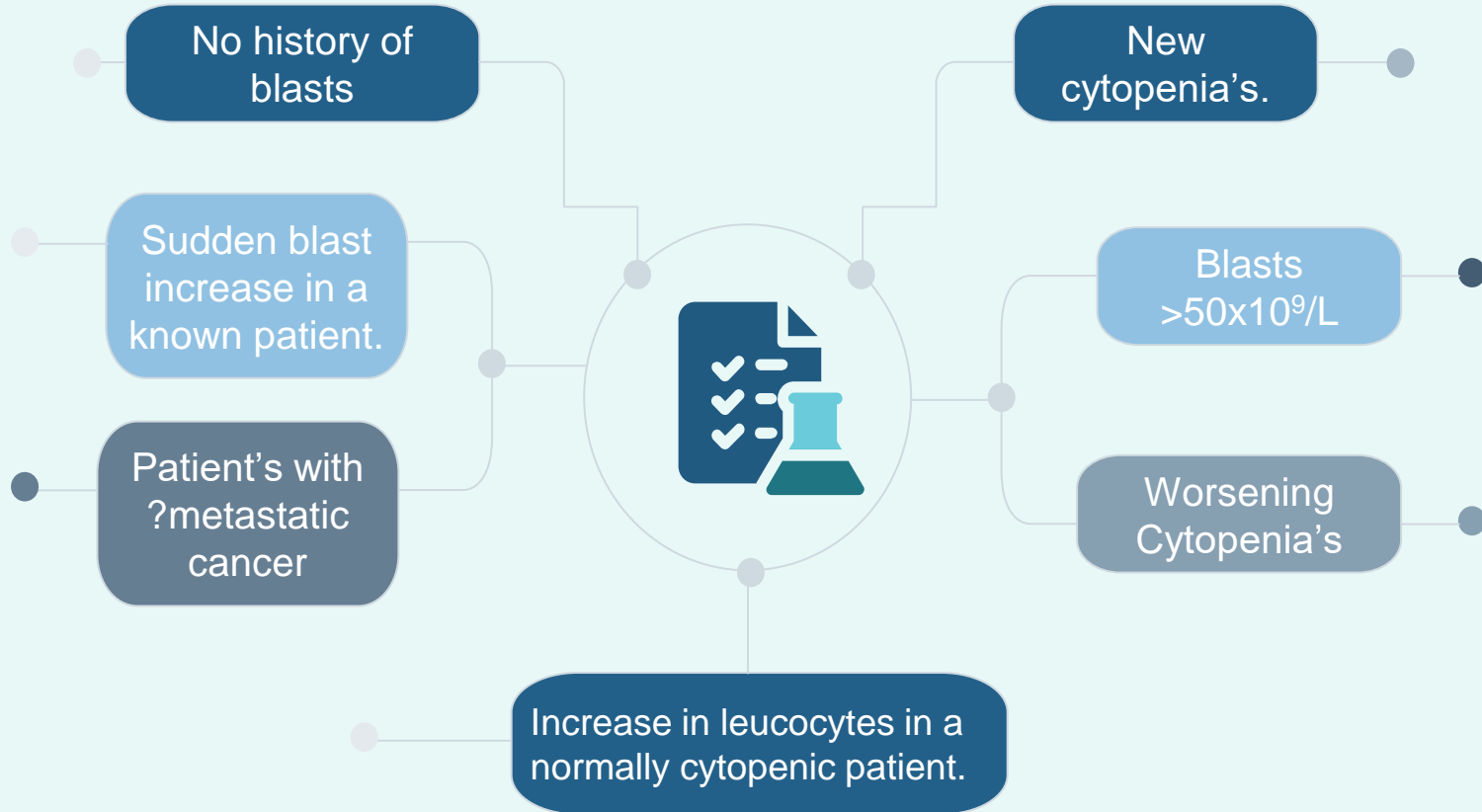
A
C

B
D

On the first when the blast count goes up.

When should films be passed to HQ?

When are blasts important?



When is it urgent?

When are blasts important?

APML is always a medical emergency- phone straight through a medic OOH!

New patient
majority

Patients with WBCC $>50 \times 10^9/L$ need to be phoned through urgently OOH.

Low blast counts can often wait if OOH, however, if in doubt- always let a medic know!

Known patient

of excess

This could indicate AML transformation- clinicians may be expecting this, but it doesn't hurt to let them know!

previously

if?

Fragments can indicate disseminated intravascular coagulation (DIC) – these should be phoned urgently!

When are blasts important?

Blast Check List!

Blast Features	How Many Blasts?	Other Cells	What's the history?	What should I do?
Granularity?	Are they the majority?	Are there any other cells?	Previous blasts?	Phone to the requester.
Vacuolation?		How mature are the other cells?	Previous cancer?	Put the film on HQ
Size?	Is this the same as previous?	Are they dysplastic?	Previous haematology diagnosis?	Contact a Haematologist.
Maturation signs?			Any known comorbidities?	Authorise.



Thanks!

Do you have any questions?

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